

(19)



Europäisches Patentamt  
European Patent Office  
Office européen des brevets



(11) Publication number:

**0 499 299 A2**

(12)

## EUROPEAN PATENT APPLICATION

(21) Application number: **92200153.2**

(51) Int. Cl.<sup>5</sup>: **A61K 47/48, A61K 9/51**

(22) Date of filing: **20.01.92**

(30) Priority: **25.01.91 US 647105**

(43) Date of publication of application:  
**19.08.92 Bulletin 92/34**

(84) Designated Contracting States:  
**AT BE CH DE DK ES FR GB GR IT LI LU MC  
NL PT SE**

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(54) **Surface modified drug nanoparticles.**

(57) Dispersible particles consisting essentially of a crystalline drug substance having a surface modifier adsorbed on the surface thereof in an amount sufficient to maintain an effective average particle size of less than about 400 nm, methods for the preparation of such particles and dispersions containing the particles. Pharmaceutical compositions containing the particles exhibit unexpected bioavailability and are useful in methods of treating mammals.

EP 0 499 299 A2

This invention relates to drug particles, methods for the preparation thereof and dispersions containing the particles. This invention further relates to the use of such particles in pharmaceutical compositions and methods of treating mammals.

Bioavailability is the degree to which a drug becomes available to the target tissue after administration. Many factors can affect bioavailability including the dosage form and various properties, e.g., dissolution rate of the drug. Poor bioavailability is a significant problem encountered in the development of pharmaceutical compositions, particularly those containing an active ingredient that is poorly soluble in water. Poorly water soluble drugs, i.e., those having a solubility less than about 10 mg/ml, tend to be eliminated from the gastrointestinal tract before being absorbed into the circulation. Moreover, poorly water soluble drugs tend to be unsafe for intravenous administration techniques, which are used primarily in conjunction with fully soluble drug substances.

It is known that the rate of dissolution of a particulate drug can increase with increasing surface area, i.e., decreasing particle size. Consequently, methods of making finely divided drugs have been studied and efforts have been made to control the size and size range of drug particles in pharmaceutical compositions. For example, dry milling techniques have been used to reduce particle size and hence influence drug absorption. However, in conventional dry milling, as discussed by Lachman et al, *The Theory and Practice of Industrial Pharmacy*, Chapter 2, "Milling", p. 45, (1986), the limit of fineness is reached in the region of 100 microns (100,000 nm) when material cakes on the milling chamber. Lachman et al note that wet grinding is beneficial in further reducing particle size, but that flocculation restricts the lower particle size limit to approximately 10 microns (10,000 nm). However, there tends to be a bias in the pharmaceutical art against wet milling due to concerns associated with contamination. Commercial airjet milling techniques have provided particles ranging in average particle size from as low as about 1 to 50  $\mu\text{m}$  (1,000 - 50,000 nm). However, such dry milling techniques can cause unacceptable levels of dust.

Other techniques for preparing pharmaceutical compositions include loading drugs into liposomes or polymers, e.g., during emulsion polymerization. However, such techniques have problems and limitations. For example, a lipid soluble drug is often required in preparing suitable liposomes. Further, unacceptably large amounts of the liposome or polymer are often required to prepare unit drug doses. Further still, techniques for preparing such pharmaceutical compositions tend to be complex. A principal technical difficulty encountered with emulsion polymerization is the removal of contaminants, such as unreacted monomer or initiator, which can be toxic, at the end of the manufacturing process.

U. S. Patent 4,540,602 (Motoyama et al) discloses a solid drug pulverized in an aqueous solution of a water-soluble high molecular substance using a wet grinding machine. Motoyama et al teach that as a result of such wet grinding, the drug is formed into finely divided particles ranging from 0.5  $\mu\text{m}$  (500 nm) or less to 5  $\mu\text{m}$  (5,000 nm) in diameter. However, there is no suggestion that particles having an average particle size of less than about 400 nm can be obtained. Attempts to reproduce the wet grinding process described by Motoyama et al resulted in particles having an average particle size much greater than 1  $\mu\text{m}$ .

EPO 275,796 describes the production of colloiddally dispersible systems comprising a substance in the form of spherical particles smaller than 500 nm. However, the method involves a precipitation effected by mixing a solution of the substance and a miscible non-solvent for the substance and results in the formation of non-crystalline nanoparticles. Furthermore, precipitation techniques for preparing particles tend to provide particles contaminated with solvents. Such solvents are often toxic and can be very difficult, if not impossible, to adequately remove to pharmaceutically acceptable levels to be practical.

U. S. Patent 4,107,288 describes particles in the size range from 10 to 1,000 nm containing a biologically or pharmacodynamically active material. However, the particles comprise a crosslinked matrix of macromolecules having the active material supported on or incorporated into the matrix.

It would be desirable to provide stable dispersible drug particles in the submicron size range which can be readily prepared and which do not appreciably flocculate or agglomerate due to interparticle attractive forces and do not require the presence of a crosslinked matrix. Moreover, it would be highly desirable to provide pharmaceutical compositions having enhanced bioavailability.

We have discovered stable, dispersible drug nanoparticles and a method for preparing such particles by wet milling in the presence of grinding media in conjunction with a surface modifier. The particles can be formulated into pharmaceutical compositions exhibiting remarkably high bioavailability.

More specifically, in accordance with this invention, there are provided particles consisting essentially of a crystalline drug substance having a surface modifier adsorbed on the surface thereof in an amount sufficient to maintain an effective average particle size of less than about 400 nm.

This invention also provides a stable dispersion consisting essentially of a liquid dispersion medium and the above-described particles dispersed therein.

In another embodiment of the invention, there is provided a method of preparing the above-described

particles comprising the steps of dispersing a drug substance in a liquid dispersion medium and applying mechanical means in the presence of grinding media to reduce the particle size of the drug substance to an effective average particle size of less than about 400 nm. The particles can be reduced in size in the presence of a surface modifier. Alternatively, the particles can be contacted with a surface modifier after attrition.

In a particularly valuable and important embodiment of the invention, there is provided a pharmaceutical composition comprising the above-described particles and a pharmaceutically acceptable carrier therefor. Such pharmaceutical composition is useful in a method of treating mammals.

It is an advantageous feature that a wide variety of surface modified drug nanoparticles free of unacceptable contamination can be prepared in accordance with this invention.

It is another advantageous feature of this invention that there is provided a simple and convenient method for preparing drug nanoparticles by wet milling in conjunction with a surface modifier, which does not result in unacceptable levels of dust as do conventional dry milling techniques.

Another particularly advantageous feature of this invention is that pharmaceutical compositions are provided exhibiting unexpectedly high bioavailability.

Still another advantageous feature of this invention is that pharmaceutical compositions containing poorly water soluble drug substances are provided which are suitable for intravenous administration techniques.

This invention is based partly on the discovery that drug particles having an extremely small effective average particle size can be prepared by wet milling in the presence of grinding media in conjunction with a surface modifier, and that such particles are stable and do not appreciably flocculate or agglomerate due to interparticle attractive forces and can be formulated into pharmaceutical compositions exhibiting unexpectedly high bioavailability. While the invention is described herein primarily in connection with its preferred utility, i.e., with respect to nanoparticulate drug substances for use in pharmaceutical compositions, it is also believed to be useful in other applications such as the formulation of particulate cosmetic compositions and the preparation of particulate dispersions for use in image and magnetic recording elements.

The particles of this invention comprise a drug substance. The drug substance exists as a discrete, crystalline phase. The crystalline phase differs from a non-crystalline or amorphous phase which results from precipitation techniques, such as described in EPO 275,796 cited above.

The invention can be practised with a wide variety of drug substances. The drug substance preferably is an organic substance present in an essentially pure form. The drug substance must be poorly soluble and dispersible in at least one liquid medium. By "poorly soluble" it is meant that the drug substance has a solubility in the liquid dispersion medium, e.g. water, of less than about 10 mg/ml, and preferably of less than about 1 mg/ml at processing temperature, e.g., room temperature. A preferred liquid dispersion medium is water. However, the invention can be practised with other liquid media in which a drug substance is poorly soluble and dispersible including, for example, aqueous salt solutions, safflower oil and solvents such as ethanol, t-butanol, hexane and glycol. The pH of the aqueous dispersion media can be adjusted by techniques known in the art.

Suitable drug substances can be selected from a variety of known classes of drugs including, for example, analgesics, anti-inflammatory agents, anthelmintics, anti-arrhythmic agents, antibiotics (including penicillins), anticoagulants, antidepressants, antidiabetic agents, antiepileptics, antihistamines, antihypertensive agents, antimuscarinic agents, antimycobacterial agents, antineoplastic agents, immunosuppressants, antithyroid agents, antiviral agents, anxiolytic sedatives (hypnotics and neuroleptics), astringents, beta-adrenoceptor blocking agents, blood products and substitutes, cardiac inotropic agents, contrast media, corticosteroids, cough suppressants (expectorants and mucolytics), diagnostic agents, diagnostic imaging agents, diuretics, dopaminergics (antiparkinsonian agents), haemostatics, immunological agents, lipid regulating agents, muscle relaxants, parasympathomimetics, parathyroid calcitonin and biphosphonates, prostaglandins, radio-pharmaceuticals, sex hormones (including steroids), anti-allergic agents, stimulants and anoretics, sympathomimetics, thyroid agents, vasodilators and xanthines. Preferred drug substances include those intended for oral administration and intravenous administration. A description of these classes of drugs and a listing of species within each class can be found in Martindale, *The Extra Pharmacopoeia*, Twenty-ninth Edition, The Pharmaceutical Press, London, 1989. The drug substances are commercially available and/or can be prepared by techniques known in the art.

Representative illustrative species of drug substances useful in the practice of this invention include:

17- $\alpha$ -pregno-2,4-dien-20-yno-[2,3-d]-isoxazol-17-ol (Danazol);  
5 $\alpha$ ,17 $\alpha$ -1'-(methylsulfonyl)-1'H-pregn-20-yno [3,2-c]-pyrazol-17-ol (Steroid A);  
[6-methoxy-4-(1-methylethyl)3-oxo-1,2-benzisothiazol-2(3H)-yl]methyl 2,6-dichlorobenzoate 1,1-dioxide (WIN 63,394);

3-amino-1,2,4-benzotriazine-1,4-dioxide (WIN 59,075);  
 piposulfam; piposulfan; camptothecin; acetaminophen; acetylsalicylic acid; amiodarone; cholestyramine;  
 colestipol; cromolyn sodium; albuterol; sucralfate; sulfasalazine; minoxidil; tempazepam; alprazolam; propox-  
 yphene; auranofin; erythromycin; cyclosporine; acyclovir; ganciclovir; etoposide; mephalan; methotrexate;  
 5 mitoxantrone; daunorubicin; doxorubicin; megestrol; tamoxifen; medroxyprogesterone; nystatin; terbutaline;  
 amphotericin B; aspirin; ibuprofen; naproxen; indomethacin; diclofenac; ketoprofen; flubiprofen; diflunisal;  
 ethyl-3,5-diacetoamido-2,4,6-triiodobenzoate (WIN 8883);  
 ethyl-(3,5-bis(acetylamino)-2,4,6-triiodobenzoyloxy)acetate (WIN 12,901); and  
 ethyl-2-(3,5-bis(acetylamino)-2,4,6-triiodobenzoyloxy)acetate (WIN 16,318).

10 In preferred embodiments of the invention, the drug substance is a steroid such as Danazol or Steroid  
 A, an antiviral agent, an anti-inflammatory agent, an antineoplastic agent, a radiopharmaceutical or a  
 diagnostic imaging agent.

The particles of this invention contain a discrete phase of a drug substance as described above having  
 a surface modifier adsorbed on the surface thereof. Useful surface modifiers are believed to include those  
 15 which physically adhere to the surface of the drug substance but do not chemically bond to the drug.

Suitable surface modifiers can preferably be selected from known organic and inorganic pharmaceutical  
 excipients. Such excipients include various polymers, low molecular weight oligomers, natural products and  
 surfactants. Preferred surface modifiers include nonionic and anionic surfactants. Representative examples  
 of excipients include gelatin, casein, lecithin (phosphatides), gum acacia, cholesterol, tragacanth, stearic  
 20 acid, benzalkonium chloride, calcium stearate, glyceryl monostearate, cetostearyl alcohol, cetomacrogol  
 emulsifying wax, sorbitan esters, polyoxyethylene alkyl ethers, e.g., macrogol ethers such as cetomacrogol  
 1000, polyoxyethylene castor oil derivatives, polyoxyethylene sorbitan fatty acid esters, e.g., the commer-  
 cially available Tweens™, polyethylene glycols, polyoxyethylene stearates, colloidal silicon dioxide, phos-  
 phates, sodium dodecylsulfate, carboxymethylcellulose calcium, carboxymethylcellulose sodium, methylcel-  
 25 lulose, hydroxyethylcellulose, hydroxypropylcellulose, hydroxypropylmethylcellulose phthalate, noncrystal-  
 line cellulose, magnesium aluminum silicate, triethanolamine, polyvinyl alcohol (PVA), and polyvinylpyr-  
 rolidone (PVP). Most of these excipients are described in detail in the *Handbook of Pharmaceutical  
 Excipients*, published jointly by the American Pharmaceutical Association and The Pharmaceutical Society  
 of Great Britain, the Pharmaceutical Press, 1986. The surface modifiers are commercially available and/or  
 30 can be prepared by techniques known in the art. Two or more surface modifiers can be used in  
 combination.

Particularly preferred surface modifiers include polyvinylpyrrolidone, tyloxapol, polaxomers, such as  
 Pluronic™ F68 and F108, which are block copolymers of ethylene oxide and propylene oxide available  
 from BASF, and poloxamines, such as Tetronic™ 908 (T908), which is a tetrafunctional block copolymer  
 35 derived from sequential addition of ethylene oxide and propylene oxide to ethylenediamine available from  
 BASF, dextran, lecithin, Aerosol OT™, which is a dioctyl ester of sodium sulfosuccinic acid, available from  
 American Cyanamid, Duponol™ P, which is a sodium lauryl sulfate, available from DuPont, Triton™ X-200,  
 which is an alkyl aryl polyether sulfonate, available from Rohm and Haas, Tween 20 and Tween 80, which  
 are polyoxyethylene sorbitan fatty acid esters, available from ICI Specialty Chemicals, Carbowax™ 3350  
 40 and 934, which are polyethylene glycols available from Union Carbide, Crodesta™ F-110, which is a  
 mixture of sucrose stearate and sucrose distearate, available from Croda Inc., Crodesta SL-40, which is  
 available from Croda Inc., and SA90HCO, which is  $C_{18}H_{37}-CH_2(CON(CH_3)CH_2(CHOH)_4CH_2OH)_2$ . Surface  
 modifiers which have found to be particularly useful include polyvinylpyrrolidone, Pluronic F-68, and lecithin.

The surface modifier is adsorbed on the surface of the drug substance in an amount sufficient to  
 45 maintain an effective average particle size of less than about 400 nm. The surface modifier does not  
 chemically react with the drug substance or itself. Furthermore, the individually adsorbed molecules of the  
 surface modifier are essentially free of intermolecular crosslinkages.

As used herein, particle size refers to a number average particle size as measured by conventional  
 particle size measuring techniques well known to those skilled in the art, such as sedimentation field flow  
 50 fractionation, photon correlation spectroscopy, or disk centrifugation. By "an effective average particle size  
 of less than about 400 nm" it is meant that at least 90% of the particles have a weight average particle size  
 of less than about 400 nm when measured by the above-noted techniques. In preferred embodiments of the  
 invention, the effective average particle size is less than about 250 nm. In some embodiments of the  
 invention, an effective average particle size of less than about 100 nm has been achieved. With reference to  
 55 the effective average particle size, it is preferred that at least 95% and, more preferably, at least 99% of the  
 particles have a particle size less than the effective average, e.g., 400 nm. In particularly preferred  
 embodiments, essentially all of the particles have a size less than 400 nm. In some embodiments,  
 essentially all of the particles have a size less than 250 nm.

The particles of this invention can be prepared in a method comprising the steps of dispersing a drug substance in a liquid dispersion medium and applying mechanical means in the presence of grinding media to reduce the particle size of the drug substance to an effective average particle size of less than about 400 nm. The particles can be reduced in size in the presence of a surface modifier. Alternatively, the particles can be contacted with a surface modifier after attrition.

A general procedure for preparing the particles of this invention is set forth below. The drug substance selected is obtained commercially and/or prepared by techniques known in the art in a conventional coarse form. It is preferred, but not essential, that the particle size of the coarse drug substance selected be less than about 100  $\mu\text{m}$  as determined by sieve analysis. If the coarse particle size of the drug substance is greater than about 100  $\mu\text{m}$ , then it is preferred that the particles of the drug substance be reduced in size to less than 100  $\mu\text{m}$  using a conventional milling method such as airjet or fragmentation milling.

The coarse drug substance selected can then be added to a liquid medium in which it is essentially insoluble to form a premix. The concentration of the drug substance in the liquid medium can vary from about 0.1 - 60%, and preferably is from 5 - 30% (w/w). It is preferred, but not essential, that the surface modifier be present in the premix. The concentration of the surface modifier can vary from about 0.1 to about 90%, and preferably is 1 - 75%, more preferably 20-60%, by weight based on the total combined weight of the drug substance and surface modifier. The apparent viscosity of the premix suspension is preferably less than about 1000 centipoise.

The premix can be used directly by subjecting it to mechanical means to reduce the average particle size in the dispersion to less than 400 nm. It is preferred that the premix be used directly when a ball mill is used for attrition. Alternatively, the drug substance and, optionally, the surface modifier, can be dispersed in the liquid medium using suitable agitation, e.g., a roller mill or a Cowles type mixer, until a homogeneous dispersion is observed in which there are no large agglomerates visible to the naked eye. It is preferred that the premix be subjected to such a premilling dispersion step when a recirculating media mill is used for attrition.

The mechanical means applied to reduce the particle size of the drug substance conveniently can take the form of a dispersion mill. Suitable dispersion mills include a ball mill, an attritor mill, a vibratory mill, and media mills such as a sand mill and a bead mill. A media mill is preferred due to the relatively shorter milling time required to provide the intended result, i.e., the desired reduction in particle size. For media milling, the apparent viscosity of the premix preferably is from about 100 to about 1000 centipoise. For ball milling, the apparent viscosity of the premix preferably is from about 1 up to about 100 centipoise. Such ranges tend to afford an optimal balance between efficient particle fragmentation and media erosion.

The grinding media for the particle size reduction step can be selected from rigid media preferably spherical or particulate in form having an average size less than about 3 mm and, more preferably, less than about 1 mm. Such media desirably can provide the particles of the invention with shorter processing times and impart less wear to the milling equipment. The selection of material for the grinding media is not believed to be critical. We have found that zirconium oxide, such as 95% ZrO stabilized with magnesia, zirconium silicate, and glass grinding media provide particles having levels of contamination which are believed to be acceptable for the preparation of pharmaceutical compositions. However, other media, such as stainless steel, titania, alumina, and 95% ZrO stabilized with yttrium, are expected to be useful. Preferred media have a density greater than about 3 g/cm<sup>3</sup>.

The attrition time can vary widely and depends primarily upon the particular mechanical means and processing conditions selected. For ball mills, processing times of up to five days or longer may be required. On the other hand, processing times of less than 1 day (residence times of one minute up to several hours) have provided the desired results using a high shear media mill.

The particles must be reduced in size at a temperature which does not significantly degrade the drug substance. Processing temperatures of less than about 30 - 40°C are ordinarily preferred. If desired, the processing equipment can be cooled with conventional cooling equipment. The method is conveniently carried out under conditions of ambient temperature and at processing pressures which are safe and effective for the milling process. For example, ambient processing pressures are typical of ball mills, attritor mills and vibratory mills. Processing pressures up to about 20 psi (1.4 kg/cm<sup>2</sup>) are typical of media milling.

The surface modifier, if it was not present in the premix, must be added to the dispersion after attrition in an amount as described for the premix above. Thereafter, the dispersion can be mixed, e.g., by shaking vigorously. Optionally, the dispersion can be subjected to a sonication step, e.g., using an ultrasonic power supply. For example, the dispersion can be subjected to ultrasonic energy having a frequency of 20 - 80 kHz for a time of about 1 to 120 seconds.

The relative amount of drug substance and surface modifier can vary widely and the optimal amount of the surface modifier can depend, for example, upon the particular drug substance and surface modifier

selected, the critical micelle concentration of the surface modifier if it forms micelles, etc. The surface modifier preferably is present in an amount of about 0.1-10 mg per square meter surface area of the drug substance. The surface modifier can be present in an amount of 0.1-90%, preferably 20-60% by weight based on the total weight of the dry particle.

5 As indicated by the following examples, not every combination of surface modifier and drug substance provides the desired results. Consequently, the applicants have developed a simple screening process whereby compatible surface modifiers and drug substances can be selected which provide stable disper-  
sions of the desired particles. First, coarse particles of a selected drug substance of interest are dispersed  
10 in a liquid in which the drug is essentially insoluble, e.g., water at 5% (w/w) and milled for 60 minutes in a DYNO-MILL under the standard milling conditions which are set forth in Example 1 which follows. The  
milled material is then divided into aliquots and surface modifiers are added at concentrations of 2, 10 and  
50% by weight based on the total combined weight of the drug substance and surface modifier. The  
15 dispersions are then sonicated (1 minute, 20 kHz) to disperse agglomerates and subjected to particle size analysis by examination under an optical microscope (1000 x magnification). If a stable dispersion is  
observed, then the process for preparing the particular drug substance surface modifier combination can be  
optimized in accordance with the teachings above. By stable it is meant that the dispersion exhibits no  
flocculation or particle agglomeration visible to the naked eye at least 15 minutes, and preferably, at least  
two days or longer after preparation.

The resulting dispersion of this invention is stable and consists of the liquid dispersion medium and the  
20 above-described particles. The dispersion of surface modified drug nanoparticles can be spray coated onto sugar spheres or onto a pharmaceutical excipient in a fluid-bed spray coater by techniques well known in the art.

Pharmaceutical compositions according to this invention include the particles described above and a  
pharmaceutically acceptable carrier therefor. Suitable pharmaceutically acceptable carriers are well known  
25 to those skilled in the art. These include non-toxic physiologically acceptable carriers, adjuvants or vehicles for parenteral injection, for oral administration in solid or liquid form, for rectal administration, and the like. A method of treating a mammal in accordance with this invention comprises the step of administering to the  
mammal in need of treatment an effective amount of the above-described pharmaceutical composition. The  
selected dosage level of the drug substance for treatment is effective to obtain a desired therapeutic  
30 response for a particular composition and method of administration. The selected dosage level therefore, depends upon the particular drug substance, the desired therapeutic effect, on the route of administration, on the desired duration of treatment and other factors. As noted, it is a particularly advantageous feature  
that the pharmaceutical compositions of this invention exhibit unexpectedly high bioavailability as illustrated  
in the examples which follow. Furthermore, it is contemplated that the drug particles of this invention  
35 provide more rapid onset of drug action in oral applications and decreased gastric irritancy.

It is contemplated that the pharmaceutical compositions of this invention will be particularly useful in  
oral and parenteral, including intravenous, administration applications. It is expected that poorly water  
soluble drug substances, which prior to this invention, could not have been administered intravenously, may  
be administered safely in accordance with this invention. Additionally, drug substances which could not  
40 have been administered orally due to poor bioavailability may be effectively administered in accordance  
with this invention.

While applicants do not wish to be bound by theoretical mechanisms, it is believed that the surface  
modifier hinders the flocculation and/or agglomeration of the particles by functioning as a mechanical or  
steric barrier between the particles, minimizing the close, interparticle approach necessary for agglomera-  
45 tion and flocculation. Alternatively, if the surface modifier has ionic groups, stabilization by electrostatic  
repulsion may result. It was surprising that stable drug particles of such a small effective average particle  
size and free of unacceptable contamination could be prepared by the method of this invention.

The following examples further illustrate the invention.

#### 50 Example 1 - PVP Modified Danazol particles prepared in a ball mill

A nanoparticulate dispersion of Danazol was prepared using a DYNO-MILL (Model KDL, manufactured  
by Willy A. Bachoffen AG Maschinenfabrik).

The following ingredients were added to a glass vessel and agitated on a roller for 24 hours to dissolve  
55 the polyvinylpyrrolidone surface modifier.

Polyvinylpyrrolidone K-15 (made by GAF) - 98 g

High purity water - 664 g

Subsequently, 327 grams of dry powdered Danazol was added to the above solution and rolled for one

week. This step aided in evenly dispersing the Danazol in the surface modifier solution, thereby reducing the treatment time required in the media mill.

The Danazol was purchased in a micronized form (average particle size of about 10 microns) from Sterling Winthrop Inc. The particles had been prepared by a conventional airjet milling technique.

5 This premix was added to a holding vessel and agitated with a conventional propeller mixer at low speed to maintain a homogeneous mixture for the media milling event. The media mill was prepared accordingly for the media milling process. The mill grinding chamber was partially filled with silica glass spheres and the premix was continuously recirculated through the media mill operating at the following conditions:

10 Grinding vessel: water jacketed stainless steel chamber

Premix flow rate: 250 ml per minute

Available volume of grinding vessel: 555 ml

Media volume: 472 ml of glass beads

Media type: size range of 0.5 - 0.75 mm silica glass beads, unleaded (distributed by Glen Mills, Inc.)

15 Recirculation time: 240 min

Residence time: 60 min

Impeller speed: 3000 RPM, tangential speed 1952 ft/min

(595 m/min)

Grinding vessel coolant: water

20 Coolant temperature: 50°F (10°C)

After recirculating the slurry for 240 minutes, a sample of the dispersion was removed and evaluated for particle size distribution using a sedimentation field flow fractionator (made by DuPont). The particles were determined to have a number average diameter of 77.5 nm and a weight average diameter of 139.6 nm. The particle size of the dispersion ranged in size from 3 - 320 nm.

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#### Example 2 - PVP modified Danazol particles prepared in a ball mill at low solids.

A nanoparticulate dispersion of Danazol was prepared using a ball mill process. A 600 ml cylindrical glass vessel (inside diameter = 3.0 inches (7.6 cm)) was filled approximately halfway with the following grinding media:

30 grinding media:

Grinding media: zirconium oxide grinding spheres (made by Zircoa, Inc.)

Media size: 0.85 - 1.18 mm diameter

Media volume: 300 ml

The following dry ingredients were added directly to this glass vessel:

35 Danazol (micronized): 10.8 g

Polyvinylpyrrolidone K-15: 3.24 g

High purity water: 201.96 g

Danzol was purchased in the micronized form (average particle size 10 microns) from Sterling Winthrop Inc. and the polyvinylpyrrolidone was K-15 grade produced by GAF.

40 The cylindrical vessel was rotated horizontally about its axis at 57% of the "critical speed". The critical speed is defined as the rotational speed of the grinding vessel when centrifuging of the grinding media occurs. At this speed the centrifugal force acting on the grinding spheres presses and holds them firmly against the inner wall of the vessel. Conditions that lead to unwanted centrifuging can be computed from simple physical principles.

45 After 5 days of ball milling, the slurry was separated from the grinding media through a screen and evaluated for particle size with the sedimentation field flow fractionator. The number average particle diameter measured was 84.9 nm and the weight average particle diameter was 169.1 nm. The particles varied in size from 26 to 340 nm. The amount and type of surface modifier was sufficient to provide colloidal stability to agglomeration and to maintain a homogeneous blend of ingredients assuring precise material delivery during subsequent processing steps.

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#### BIOAVAILABILITY TESTING

55 Bioavailability of Danazol from the nanoparticulate dispersion described above was compared to that from a suspension of unmilled Danazol in fasted male beagle dogs. The unmilled material was prepared as a suspension in the same manner as the dispersion, with the exception of the ball milling process. Both formulations were administered to each of five dogs by oral gavage and plasma obtained via a cannula in the cephalic vein. Plasma Danazol levels were monitored over 24 hours. The relative bioavailability of

Danazol from the nanoparticulate dispersion was 15.9 fold higher than from the Danazol suspension containing Danazol particles having an average particle size of about 10 microns prepared by conventional airjet milling. Comparison of oral plasma levels with dose corrected plasma levels following intravenous administration of Danazol gave a mean absolute bioavailability ( $\pm$  SEM) of  $82.3 \pm 10.1\%$  for the nanoparticulate dispersion and  $5.1 \pm 1.9\%$  for the unmilled material.

#### Example 3 - PVP modified Danazol particles prepared in a ball mill at high solids

A nanoparticle dispersion of Danazol was prepared using 1 mm diameter glass grinding media (.85 - 1.18 mm from Potters Industries). A cylindrical glass vessel having a diameter of 2.75 inches (7.0 cm) with a volume of 400 ml was charged with 212 ml of unleaded glass grinding media. The following ingredients were added to this vessel:

30.4 g of micronized Danazol

9.12 g of Polyvinylpyrrolidone K-15

112.48 g of high purity water

This vessel was rotated horizontally on its axis at a controlled rotational speed of 80.4 revolutions per minute (50% of critical speed) for 5 days. The slurry was immediately separated from the grinding media and evaluated for particle size and grinding media attrition using inductively coupled plasma emissions (ICP). The particle size measured with a sedimentation field flow fractionator yielded a number average diameter of 112.7 nm and a weight average diameter of 179.3 nm. The extent of media attrition was measured to establish the purity of the final dispersion using an inductively coupled plasma-atomic emission spectroscopy method. The level of silicon in the final dispersion was less than 10 parts of elemental silicon per million parts of the slurry.

#### Example 4 - PVP modified Danazol particles

A nanoparticle dispersion of Danazol was prepared for clinical evaluation using a ball milling dispersion method. This dispersion was prepared by milling with glass grinding media. The grinding media used was: Media type: 0.85 - 1.18 mm unleaded glass spheres

Media quantity: 6100 ml

The media was added to a 3 gallon porcelain jar. The following ingredients were then added to the jar:

1000 g Danazol (micronized)

300 g Polyvinylpyrrolidone K-15

3700 g high purity water

The vessel was rolled 5 days at a rotational speed of 39.5 revolutions per minute (50% critical speed). The liquid slurry was separated from the grinding media with a screen and used to prepare solid oral doses for clinical studies. The dispersion was assessed for particle size using the sedimentation field flow fractionator and was measured to have a number average diameter of 134.9 nm and a weight average diameter of 222.2 nm. The level of contamination from the grinding media was measured (by ICP) to be 36 parts of silicon per million parts of dispersion. Less than 5 ppm of aluminum was detected. X-ray powder diffraction data of the starting powder was compared with the dispersed Danazol and showed the crystal structure morphology of the solid dispersed particles was unchanged by the dispersion process.

#### Example 5 - PVP modified Danazol particles

A nanoparticulate dispersion of Danazol was prepared using a laboratory media mill and glass grinding media. The media mill was equipped with a 50 ml grinding chamber and the mill was a "Mini" Motormill manufactured by Eiger Machinery Inc.

The media mill was operated at the following process conditions:

Bead charge: 42.5 ml glass spheres

Rotor speed: 5000 RPM (2617 feet per minute (798 m/min) tangential speed)

Grinding media: 0.75 - 1.0 mm unleaded glass beads (distributed by Glens Mills)

The dispersion formula was prepared by dissolving 27 g of polyvinylpyrrolidone in 183 g of water and agitated in a steel vessel with a 50 mm "Cowles" type blade until the solution was clear and free of undissolved PVP polymer. The rotational speed of the mixer was maintained at 5000 RPM. 90 g of micronized Danazol was slowly added to this blend with the same mixing for 30 min. 200 cc of the premix was added to the holding tank of the mill and recirculated for 5 hours and 51 minutes. The final residence time in the grinding zone was 40 minutes.



The final average particle size was measured and determined to have a number average diameter of 79.9 nm and a weight average diameter of 161.2 nm. The particles varied in size from 30 - 415 nm. The level of attrition from erosion of the grinding media and grinding vessel were measured (by ICP) to be 170 ppm of iron and 71 ppm silicon. The crystal structure was determined by X-ray diffraction to be unchanged by the dispersion process.

#### Example 6 - Lecithin modified Steroid A particles

A nanoparticulate dispersion of Steroid A was prepared by ball milling with zirconium oxide grinding beads. The dispersion was prepared in the absence of a surface modifier and a post addition of Lecithin and a sonication step were required to stabilize the dispersed phase of Steroid A and prevent agglomeration and rapid sedimentation.

A fine particle dispersion of Steroid A was prepared by ball milling the following ingredients:

- 5 g Steroid A
- 95 g high purity water

Steroid A was in the form of unmilled coarse grains having a particle size of about 100  $\mu\text{m}$  and ranging in size up to about 400  $\mu\text{m}$ .

The following process conditions were used:

- Media: 135 ml
- Vessel volume: 240 ml
- Media type: 0.85 - 1.18 mm Zirbeads (manufactured by Zircoa Inc.)
- Milling time: 4 days
- Milling speed: 86 RPM (50% critical speed)

After four days of ball milling the slurry was separated from the grinding media through a screen. One gram of this unstabilized slurry was added to 10 g of an aqueous solution of Lecithin (1% Centrox "P" by weight in high purity water, Lecithin manufactured by Central Soya Company, Inc.) and mixed by vigorous shaking, followed by a sonication step for 20 seconds using an ultrasonic horn (Model 350 Branson Ultrasonic Power Supply, Horn Diameter = 0.5 inch (1.27 cm), Power setting = 2). The slurry was sized under a microscope. An Olympus BH-2 optical microscope equipped with phase contrast illumination was used to observe the size and condition of the dispersion.

A drop of the above dilute slurry was placed between a microscope slide and glass cover slip and observed microscopically at high magnification (1,000 times) and compared to the slurry similarly diluted with water only (no surface modifier). The unmodified dispersion exhibited extensive particle agglomeration. The particle size of the unmodified dispersion was more than 10 microns and the unmodified dispersion exhibited no Brownian Motion. Brownian motion is the oscillatory or jiggling motion exhibited by particles in a liquid that fall in the size range of less than about 1 micron. The Lecithin modified particles exhibited rapid Brownian motion. The thus observed dispersion had the characteristics and appearance consistent with a number average particle size of less than 400 nm. Furthermore, it is expected that additional milling would lead to further particle size reduction.

#### Example 7 - Alkyl aryl polyether sulfonate modified Steroid A

Example 6 was repeated except that the Lecithin was replaced with Triton X-200 (manufactured by Rohm and Haas). Similar results were observed.

#### Example 8 - Gum acacia modified Steroid A

Example 6 was repeated except that the Lecithin was replaced with gum acacia (available from Eastman Kodak Co.) Similar results were observed.

#### Example 9 - Sodium lauryl sulfate modified Steroid A

Example 6 was repeated except that the Lecithin was replaced with sodium lauryl sulfate (available as Duponol ME from DuPont, Inc.). Similar results were observed.

#### Example 10 - Steroid A modified with a dioctylester of sodium sulfosuccinic acid

Example 6 was repeated except that the Lecithin was replaced with Aerosol OT (available from American Cyanamid Chemical Products, Inc.). Similar results were observed.

Example 11 - Steroid A modified with a block copolymer of ethylene oxide and propylene oxide

Example 6 was repeated except that the Lecithin was replaced with Pluronic F68 (available from BASF Corp.). Similar results were observed.

Example 12 - Steroid A modified with a block copolymer of ethylene oxide and propylene oxide

A nanoparticulate dispersion of Steroid A was prepared by ball milling with zirconium oxide grinding media for 5 days. 70 cc of grinding media were added to a 115 cc vessel followed by:

2.5 g Steroid A

0.75 g of Pluronic F68

46.75 g high purity water

The resulting mixture was ball milled for 5 days at 50% of the critical rotational speed. The final dispersion was separated from the grinding media and microscopically evaluated for particle size as in Example 6. The dispersion exhibited rapid Brownian Motion and no particles were larger than 1 micron. Most particles were less than 400 nm.

Example 13 - Lecithin modified Steroid A particles

Example 12 was repeated except that the Pluronic F68 was replaced with Centrox P. No particles larger than 1 micron were observed microscopically and most were less than 400 nm.

Example 14 - Steroid A particles modified with a block copolymer of ethylene oxide and propylene oxide

A nanoparticulate dispersion of Steroid A was prepared by a ball milling process. The following ingredients were added to a cylindrical 0.95 l vessel. The vessel was filled approximately halfway with the following grinding media:

Grinding media: 0.85 - 1.18 mm diameter zirconium oxide spheres (made by Zircoa)

The following dispersion ingredients were added directly to the glass vessel:

18 g Steroid A

4.5 g Pluronic F68 (purchased from BASF Corp.)

336.6 g high purity water

Steroid A was purchased from Sterling Winthrop Inc. in the form of unmilled tabular crystals having an average particle size of approximately 100  $\mu$ m.

The vessel was rotated concentrically on its axis at 50% critical speed for 5 days. After this time 4.45 g of Pluronic F68 was added to the slurry and rolled for 5 more days at the same conditions. The slurry was then discharged and separated from the grinding media and evaluated for particle size using the sedimentation field flow fractionator. The number average particle size measured was 204.6 nm and the weight average particle size was 310.6 nm. The particle size distribution ranged from approximately 68 - 520 nm. The dispersion was examined with an optical microscope. It exhibited excellent particle integrity, free of flocculation and agglomeration. The dispersion particles exhibited rapid Brownian motion.

BIOAVAILABILITY TESTING

Bioavailability of Steroid A from the nanoparticulate dispersion described above was compared to that from a suspension of unmilled Steroid A (having an average particle size of about 100  $\mu$ m) in male beagle dogs. The unmilled material was prepared as a suspension in the same manner as the dispersion, with the exception of the ball milling process. Both formulations were administered to each of five dogs by oral gavage and plasma obtained via a cannula in the cephalic vein. Plasma Steroid A levels were monitored over 24 hours. The relative bioavailability of Steroid A from the nanoparticulate dispersion was 7.1 fold higher than from the unmilled Steroid A suspension. Comparison of oral plasma levels with dose corrected plasma levels following intravenous administration of Steroid A gave a mean absolute bioavailability ( $\pm$  SEM) of  $14.8 \pm 3.5\%$  for the nanoparticulate dispersion and  $2.1 \pm 1.0\%$  for the unmilled material.

Comparative Example A

A dispersion of Steroid A was prepared using a ball milling process with zirconium oxide grinding beads. The dispersion was prepared in the absence of a surface modifier and a post-sonication step was used to minimize flocculation and reaggregation.

A fine particle dispersion was prepared by ball milling the following ingredients:

5 5 g Steroid A

95 g high purity water

The following process conditions were used:

Grinding media: 135 ml

Vessel volume: 240 ml

10 Grinding media: 0.85 - 1.18 mm Zirbeads XR

Milling time: 4 days

Milling speed: 86 RPM (50% critical speed)

After four days of ball milling, the slurry was separated from the grinding media through a screen. One gram of the unstablized slurry was blended with 10 grams of high purity water and mixed by vigorous shaking, followed by a sonication step for 20 seconds using an ultrasonic horn (Model 350 Branson Ultrasonic Power Supply, Horn diameter = 0.5 inch, Power setting = 2). The slurry was sized under a microscope. An optical microscope equipped with phase contrast illumination was used to observe the condition of the dispersion.

20 A drop of the dilute slurry was placed between a microscope slide and a glass cover slip and observed at high magnification (400X). The dispersion exhibited severe particle aggregation. The aggregate size was greater than 10 microns and exhibited no Brownian particle movement.

#### Examples 15-49

25 Table 1 is a summary of additional examples of the invention. Each of the examples in Table 1 resulted in particles having an effective average particle size of less than 400 nm.

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TABLE 1

	Example	Class	Drug Substance	Surface Modifier	Particle Size
5	15.	anti-inflammatory	5% naproxen	5% PVP	250 nm
	16.	anti-inflammatory	5% naproxen	3% F68	267 nm
	17.	anti-inflammatory	5% indomethacin	1% F68	228 nm
10	18.	anti-inflammatory	5% indomethacin	1% PVA	331 nm
	19.	anti-inflammatory	5% indomethacin	1% PVP	216 nm
	20.	anti-inflammatory	5% indomethacin	1% F108	235 nm
15	21.	anti-inflammatory	3% WIN 63,394	0.5% F68	262 nm
	22.	anti-inflammatory	4% WIN 63,394	3% F68	255 nm
	23.	anti-inflammatory	3% WIN 63,394	10% F68	231 nm
	24.	antineoplastic	1% etoposide	1% Crodesta F-110	-300 nm
20	25.	antineoplastic	1% etoposide	1% Crodesta SL-40	-300 nm
25	26.	antineoplastic	1% etoposide	1% F68	-300 nm
	27.	antineoplastic	1% etoposide	1% F108	-300 nm
	28.	antineoplastic	1% etoposide	1% gum acacia	-300 nm
	29.	antineoplastic	1% etoposide	1% PVA	-300 nm
30	30.	antineoplastic	1% camptothecin	0.6% gum acacia	298 nm
	31.	antineoplastic	1% camptothecin	1.1% PVA	236 nm
	32.	antineoplastic	1% camptothecin	1% T908	256 nm
35	33.	antineoplastic	5% piposulfam	1.25% Crodesta F-110	-300 nm
	34.	antineoplastic	5% piposulfam	1.25% gum acacia	-300 nm
	35.	antineoplastic	5% piposulfam	5% PVA	320 nm
40	36.	radiopharmaceutical	2.5% WIN 59,075	3% PVP	359 nm

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Example	Class	Drug	Surface	Particle
		Substance	Modifier	Size
5	37. diagnostic imaging agent	10% WIN 8883	2% T908	166 nm
	38. diagnostic imaging agent	20% WIN 8883	3.3% T908	180 nm
10	39. diagnostic imaging agent	20% WIN 8883	3.3% T908 (isotonic phosphate buffered saline, pH= 7.4)	159 nm
15	40. diagnostic imaging agent	20% WIN 8883	3.3% T908 (0.1M phosphate buffer pH=7.5)	167 nm
20	41. diagnostic imaging agent	10% WIN 8883	1% SA90HCO 1% Tween 20	194 nm
	42. diagnostic imaging agent	10% WIN 8883	1% SA90HCO	193 nm
25	43. diagnostic imaging agent	40% WIN 8883	3.3% T908	329 nm
30	44. diagnostic imaging agent	10% WIN 8883	2% Tween 20	241 nm
	45. diagnostic imaging agent	10% WIN 12,901	2% T908	238 nm
35	46. diagnostic imaging agent	20% WIN 12,901	3.3% T908 (phosphate buffer, pH=6.5)	289 nm
40	47. diagnostic imaging agent	10% WIN 16,318	2% Tween 80	219 nm
	48. anti-inflammatory	3% ibuprofen	2% F68	-250 nm
45	49. anti-inflammatory	3% ibuprofen	2% F68 (in 0.1M HCl)	-375 nm

These examples demonstrate that the wet grinding process of this invention is broadly applicable to a wide variety of classes of poorly-soluble drug substances including steroids, anti-inflammatory agents, antineoplastic agents, radiopharmaceutical agents and diagnostic imaging agents having radically different chemical structures. Additionally, these examples demonstrate that the invention can be practiced in conjunction with a variety of surface modifiers and at a variety of surface modifier concentrations.

Furthermore, laboratory work has demonstrated that particles prepared according to this invention have exhibited a variety of unexpected properties, particularly with respect to increased bioavailability. For example, as described above, pharmaceutical compositions containing Steroid A and Danazol according to this invention have unexpectedly exhibited 7 and 16 fold increases in bioavailability compared to dispersions prepared by conventional techniques. Aqueous dispersions of WIN 63,394 prepared according to this invention resulted in an increase in bioavailability of 37-fold when compared to a conventional dispersion of

WIN 63,394. The dispersions were administered at a dose of 5mg WIN 63,394 per kilogram of body weight to three dogs in the fasted state as a two way crossover study. Serial blood samples were withdrawn and analyzed by HPLC for WIN 63,394 concentrations. The relative bioavailabilities were calculated from the area under the curve for concentration versus time plots. Such increased bioavailability is particularly advantageous inasmuch as drug substances in the form of the particles of the instant invention can achieve the same therapeutic effect as substantially greater dosages of drug substances prepared by prior art techniques.

In addition, pharmaceutical compositions containing particles of this invention have exhibited improved dose proportionality and decreased fed-fasted variability. Further, particles of the invention comprising naproxen or indomethacin, when administered orally, have resulted in more rapid onset of action compared to conventional naproxen and indomethacin formulations. Moreover, certain of the particles of the invention have been found to be extraordinarily useful in x-ray contrast compositions.

### Claims

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1. Particles consisting essentially of a crystalline drug substance having a surface modifier adsorbed on the surface thereof in an amount sufficient to maintain an effective average particle size of less than about 400 nm.

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2. The particles of claim 1 having an effective average particle size of less than 250 nm.

3. The particles of claim 1 having an effective average particle size of less than 100 nm.

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4. The particles of claim 1 wherein said drug substance is selected from analgesics, anti-inflammatory agents, anthelmintics, anti-arrhythmic agents, antibiotics, anticoagulants, antidepressants, antidiabetic agents, antiepileptics, antihistamines, antihypertensive agents, antimuscarinic agents, antimycobacterial agents, antineoplastic agents, immunosuppressants, antithyroid agents, antiviral agents, anxiolytic sedatives, astringents, beta-adrenoceptor blocking agents, blood products and substitutes, cardiac suppressants agents, contrast media, corticosteroids, cough suppressants, diagnostic agents, diagnostic imaging agents, diuretics, dopaminergics, haemostatics, immunological agents, lipid regulating agents, muscle relaxants, parasympathomimetics, parathyroid calcitonin and biphosphonates, prostaglandins, radio-pharmaceuticals, sex hormones, steroids, anti-allergic agents, stimulants and anoretics, sympathomimetics, thyroid agents, vasodilators and xanthines.

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5. The particles of claim 1 wherein said drug substance is selected from an antiviral agent, an anti-inflammatory agent, an antineoplastic agent, a radiopharmaceutical agent, and a diagnostic imaging agent.

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6. The particles of claim 1 wherein said drug substance is selected from the group consisting of Danazol, 5 $\alpha$ ,17 $\alpha$ ,1'-1'-(methylsulfonyl)-1'H-pregn-20-yno-[3,2-c]-pyrazol-17-ol, pipsulfam, pipsulfan, campothecin, and ethyl-3,5-diacetamido-2,4,6-triodobenzoate.

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7. The particles of claim 1 wherein said surface modifier is selected from the group consisting of gelatin, casein, lecithin, gum acacia, cholesterol, tragacanth, stearic acid, benzalkonium chloride, calcium stearate, glyceryl monostearate, cetostearyl alcohol, cetomacrogol emulsifying wax, sorbitan esters, polyoxyethylene alkyl ethers, polyoxyethylene castor oil derivatives, polyoxyethylene sorbitan fatty acid esters, polyethylene glycols, polyoxyethylene stearates, colloidol silicon dioxide, phosphates, sodium dodecylsulfate, carboxymethylcellulose calcium, carboxymethylcellulose sodium, methylcellulose, hydroxyethylcellulose, hydroxypropylcellulose, hydroxypropylmethylcellulose phthalate, noncrystalline cellulose, magnesium aluminum silicate, triethanolamine, polyvinyl alcohol, polyvinylpyrrolidone, an ethylene oxide-propylene oxide block copolymer, an alkyl aryl polyether sulfonate, and a dioctylester of sodium sulfosuccinic acid.

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8. The particles of claim 1 wherein said surface modifier is present in an amount of 0.1 to 90% by weight based on the total weight of the dry particle.

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9. Particles according to claim 1 consisting essentially of crystalline Danazol having polyvinyl pyrrolidone adsorbed on the surface thereof in an amount sufficient to maintain an effective average particle size of

less than 100 nm.

10. Particles according to claim 1 consisting essentially of crystalline  $5\alpha,17\alpha,1'$ -(methylsulfonyl)-1'H-pregn-20-yno-[3,2-c]pyrazol-17-ol having an ethylene oxide propylene-oxide block copolymer adsorbed on the surface thereof in an amount sufficient to maintain an effective average particle size of less than about 400 nm.
11. A stable dispersion consisting essentially of a liquid dispersion medium and the particles of any one of claims 1-10.
12. The dispersion of claim 11 wherein said dispersion medium is water.
13. The dispersion of claim 11 wherein said dispersion medium is selected from the group consisting of safflower oil, ethanol, t-butanol, hexane and glycol.
14. A pharmaceutical composition comprising the particles of any one of claims 1-10 and a pharmaceutically acceptable carrier therefor.
15. A method of treating a mammal comprising the step of administering to the mammal an effective amount of the pharmaceutical composition of claim 14.
16. A method of preparing the particles of claim 1 comprising the steps of dispersing a drug substance in a liquid dispersion medium and wet grinding said drug substance in the presence of rigid grinding media having an average particle size of less than 3 mm and a surface modifier to reduce the particle size of said drug substance to an effective average particle size of less than about 400 nm.
17. A method of preparing the particles of claim 1 comprising the steps of dispersing a drug substance in a liquid dispersion medium, wet grinding said drug substance in the presence of rigid grinding media having an average particle size of less than 3 mm, and thereafter contacting said drug substance with a surface modifier by mixing said surface modifier with said dispersion medium to form particles having an effective average particle size of less than about 400 nm.
18. The method of claim 18 further including the step of subjecting the dispersion medium containing said drug substance and said surface modifier to ultrasonic energy.
19. The method of claim 17 or 18 wherein said grinding media have a density greater than 3 g/cm<sup>3</sup>.
20. The method of claim 17 or 18 wherein said grinding media have an average particle size of less than 1 mm.

(19)



Europäisches Patentamt  
European Patent Office  
Office européen des brevets



(11) Publication number:

**0 499 299 A3**

(12)

**EUROPEAN PATENT APPLICATION**(21) Application number: **92200153.2**(51) Int. Cl.<sup>5</sup>: **A61K 9/51, A61K 49/04,  
B01J 13/00**(22) Date of filing: **20.01.92**(30) Priority: **25.01.91 US 647105**(43) Date of publication of application:  
**19.08.92 Bulletin 92/34**(84) Designated Contracting States:  
**AT BE CH DE DK ES FR GB GR IT LI LU MC  
NL PT SE**(86) Date of deferred publication of the search report:  
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**Harrow, Middlesex HA1 4TY (GB)**(54) **Surface modified drug nanoparticles.**

(57) Dispersible particles consisting essentially of a crystalline drug substance having a surface modifier adsorbed on the surface thereof in an amount sufficient to maintain an effective average particle size of less than about 400 nm, methods for the preparation of such particles and dispersions containing the particles. Pharmaceutical compositions containing the particles exhibit unexpected bioavailability and are useful in methods of treating mammals.

**EP 0 499 299 A3**







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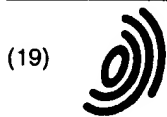
# PARTIAL EUROPEAN SEARCH REPORT

Application Number

EP 92 20 0153

Page 2

DOCUMENTS CONSIDERED TO BE RELEVANT			CLASSIFICATION OF THE APPLICATION (Int. Cl. 5)
Category	Citation of document with indication, where appropriate, of relevant passages	Relevant to claim	
X	WO-A-8 400 294 (SCHRÖDER U.) February 284 * page 1, line 21 - line 29 * * page 3, line 15 - line 20; claims * * page 7, line 18 - line 26 * * page 8, line 17 - line 23 * ---	1-20	
X	WO-A-9 015 593 (YTKEMISKA INSTITUTET) 27 December 1990 * page 1, line 2 - line 4 * * page 3, line 18 - line 34 * * page 4 - page 6 * * page 7, line 17 - page 8, line 12 * * page 9, line 24 - line 28; claims *	1-20	
Y	---	1-20	TECHNICAL FIELDS SEARCHED (Int. Cl. 5)
Y	US-A-3 192 118 (O. A. BATTISTA) 29 June 1965 * claims *	1-20	
P,A	WO-A-9 115 193 (RHONE-POULENC RORER S.A.) 17 October 1991 * page 1, line 28 - page 2, line 10 * * page 2, line 29 - page 3, line 34; claims *	1-20	
D,X	DE-A-3 013 839 (FREUND INDUSTRIAL CO.) 30 October 1980 * claims; examples 6,8,13,13 * & US-A-4 540 602 -----	1-20	



(19)

Europäisches Patentamt

European Patent Office

Office européen des brevets



(11)

**EP 0 499 299 B1**

(12)

## EUROPEAN PATENT SPECIFICATION

(45) Date of publication and mention  
of the grant of the patent:  
**16.08.2000 Bulletin 2000/33**

(51) Int. Cl.<sup>7</sup>: **A61K 9/51, A61K 49/04,  
B01J 13/00**

(21) Application number: **92200153.2**

(22) Date of filing: **20.01.1992**

### (54) **Surface modified drug nanoparticles**

Oberflächenmodifizierte Arzneimittel-Nanopartikel

Nanoparticules de médicaments à surface modifiée

(84) Designated Contracting States:  
**AT BE CH DE DK ES FR GB GR IT LI LU MC NL  
PT SE**

(30) Priority: **25.01.1991 US 647105**

(43) Date of publication of application:  
**19.08.1992 Bulletin 1992/34**

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(56) References cited:  
**WO-A-84/00294**                      **WO-A-90/15593**  
**WO-A-91/15193**                      **DE-A- 3 013 839**  
**DE-A- 3 722 837**                      **FR-A- 2 304 326**  
**US-A- 3 192 118**

• **CHEMICAL ABSTRACTS, vol. 111, no. 8, 1989,**  
**Columbus, Ohio, US; abstract no. 63850,**

#### Remarks:

The file contains technical information submitted  
after the application was filed and not included in  
this specification

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**EP 0 499 299 B1**

## Description

[0001] This invention relates to drug particles, methods for the preparation thereof and dispersions containing the particles. This invention further relates to the use of such particles in pharmaceutical compositions and methods of treating mammals.

[0002] Bioavailability is the degree to which a drug becomes available to the target tissue after administration. Many factors can affect bioavailability including the dosage form and various properties, e.g., dissolution rate of the drug. Poor bioavailability is a significant problem encountered in the development of pharmaceutical compositions, particularly those containing an active ingredient that is poorly soluble in water. Poorly water soluble drugs, i.e., those having a solubility less than about 10 mg/ml, tend to be eliminated from the gastrointestinal tract before being absorbed into the circulation. Moreover, poorly water soluble drugs tend to be unsafe for intravenous administration techniques, which are used primarily in conjunction with fully soluble drug substances.

[0003] It is known that the rate of dissolution of a particulate drug can increase with increasing surface area, i.e., decreasing particle size. Consequently, methods of making finely divided drugs have been studied and efforts have been made to control the size and size range of drug particles in pharmaceutical compositions. For example, dry milling techniques have been used to reduce particle size and hence influence drug absorption. However, in conventional dry milling, as discussed by Lachman *et al*, *The Theory and Practice of Industrial Pharmacy*, Chapter 2, "Milling", p. 45, (1986), the limit of fineness is reached in the region of 100 microns (100,000 nm) when material cakes on the milling chamber. Lachman *et al* note that wet grinding is beneficial in further reducing particle size, but that flocculation restricts the lower particle size limit to approximately 10 microns (10,000 nm). However, there tends to be a bias in the pharmaceutical art against wet milling due to concerns associated with contamination. Commercial airjet milling techniques have provided particles ranging in average particle size from as low as about 1 to 50  $\mu\text{m}$  (1,000 - 50,000 nm). However, such dry milling techniques can cause unacceptable levels of dust.

[0004] Other techniques for preparing pharmaceutical compositions include loading drugs into liposomes or polymers, e.g., during emulsion polymerization. However, such techniques have problems and limitations. For example, a lipid soluble drug is often required in preparing suitable liposomes. Further, unacceptably large amounts of the liposome or polymer are often required to prepare unit drug doses. Further still, techniques for preparing such pharmaceutical compositions tend to be complex. A principal technical difficulty encountered with emulsion polymerization is the removal of contaminants, such as unreacted monomer or initiator, which can be toxic, at the end of the manufacturing process.

[0005] U. S. Patent 4,540,602 (Motoyama *et al*) discloses a solid drug pulverized in an aqueous solution of a water-soluble high molecular substance using a wet grinding machine. Motoyama *et al* teach that as a result of such wet grinding, the drug is formed into finely divided particles ranging from 0.5  $\mu\text{m}$  (500 nm) or less to 5  $\mu\text{m}$  (5,000 nm) in diameter. However, there is no suggestion that particles having an average particle size of less than 400 nm can be obtained. Attempts to reproduce the wet grinding process described by Motoyama *et al* resulted in particles having an average particle size much greater than 1  $\mu\text{m}$ .

[0006] EPO 275,796 describes the production of colloiddally dispersible systems comprising a substance in the form of spherical particles smaller than 500 nm. However, the method involves a precipitation effected by mixing a solution of the substance and a miscible non-solvent for the substance and results in the formation of non-crystalline nanoparticles. Furthermore, precipitation techniques for preparing particles tend to provide particles contaminated with solvents. Such solvents are often toxic and can be very difficult, if not impossible, to adequately remove to pharmaceutically acceptable levels to be practical.

[0007] U. S. Patent 4,107,288 describes particles in the size range from 10 to 1,000 nm containing a biologically or pharmacodynamically active material. However, the particles comprise a crosslinked matrix of macromolecules having the active material supported on or incorporated into the matrix.

[0008] It would be desirable to provide stable dispersible drug particles in the submicron size range which can be readily prepared and which do not appreciably flocculate or agglomerate due to interparticle attractive forces and do not require the presence of a crosslinked matrix. Moreover, it would be highly desirable to provide pharmaceutical compositions having enhanced bioavailability.

[0009] We have discovered stable, dispersible drug nanoparticles and a method for preparing such particles by wet milling in the presence of grinding media in conjunction with a surface modifier. The particles can be formulated into pharmaceutical compositions exhibiting remarkably high bioavailability.

[0010] More specifically, in accordance with this invention, there are provided particles consisting essentially of a crystalline drug substance having a surface modifier adsorbed on the surface thereof in an amount sufficient to maintain an effective average particle size of less than 400 nm.

[0011] This invention also provides a stable dispersion consisting essentially of a liquid dispersion medium and the above-described particles dispersed therein.

[0012] In another embodiment of the invention, there is provided a method of preparing the above-described parti-

cles comprising the steps of dispersing a drug substance in a liquid dispersion medium and applying mechanical means in the presence of grinding media to reduce the particle size of the drug substance to an effective average particle size of less than 400 nm. The particles can be reduced in size in the presence of a surface modifier. Alternatively, the particles can be contacted with a surface modifier after attrition.

5 [0013] In a particularly valuable and important embodiment of the invention, there is provided a pharmaceutical composition comprising the above-described particles and a pharmaceutically acceptable carrier therefor. Such pharmaceutical composition is useful in a method of treating mammals.

[0014] It is an advantageous feature that a wide variety of surface modified drug nanoparticles free of unacceptable contamination can be prepared in accordance with this invention.

10 [0015] It is another advantageous feature of this invention that there is provided a simple and convenient method for preparing drug nanoparticles by wet milling in conjunction with a surface modifier, which does not result in unacceptable levels of dust as do conventional dry milling techniques.

[0016] Another particularly advantageous feature of this invention is that pharmaceutical compositions are provided exhibiting unexpectedly high bioavailability.

15 [0017] Still another advantageous feature of this invention is that pharmaceutical compositions containing poorly water soluble drug substances are provided which are suitable for intravenous administration techniques.

[0018] This invention is based partly on the discovery that drug particles having an extremely small effective average particle size can be prepared by wet milling in the presence of grinding media in conjunction with a surface modifier, and that such particles are stable and do not appreciably flocculate or agglomerate due to interparticle attractive forces and can be formulated into pharmaceutical compositions exhibiting unexpectedly high bioavailability.

20 [0019] The particles of this invention comprise a drug substance. The drug substance exists as a discrete, crystalline phase. The crystalline phase differs from a non-crystalline or amorphous phase which results from precipitation techniques, such as described in EPO 275,796 cited above.

[0020] The invention can be practised with a wide variety of drug substances. The drug substance preferably is an organic substance present in an essentially pure form. The drug substance must be poorly soluble and dispersible in at least one liquid medium. By "poorly soluble" it is meant that the drug substance has a solubility in the liquid dispersion medium, e.g. water, of less than about 10 mg/ml, and preferably of less than about 1 mg/ml at processing temperature, e.g., room temperature. A preferred liquid dispersion medium is water. However, the invention can be practised with other liquid media in which a drug substance is poorly soluble and dispersible including, for example, aqueous salt solutions, safflower oil and solvents such as ethanol, t-butanol, hexane and glycol. The pH of the aqueous dispersion media can be adjusted by techniques known in the art.

30 [0021] Suitable drug substances can be selected from a variety of known classes of drugs including, for example, analgesics, anti-inflammatory agents, anthelmintics, anti-arrhythmic agents, antibiotics (including penicillins), anticoagulants, antidepressants, antidiabetic agents, antiepileptics, antihistamines, antihypertensive agents, antimuscarinic agents, antimycobacterial agents, antineoplastic agents, immunosuppressants, antithyroid agents, antiviral agents, anxiolytic sedatives (hypnotics and neuroleptics), astringents, beta-adrenoceptor blocking agents, blood products and substitutes, cardiac inotropic agents, contrast media, corticosteroids, cough suppressants (expectorants and mucolytics), diagnostic agents, diagnostic imaging agents, diuretics, dopaminergics (antiparkinsonian agents), haemostatics, immunological agents, lipid regulating agents, muscle relaxants, parasympathomimetics, parathyroid calcitonin and

40 biphosphonates, prostaglandins, radio-pharmaceuticals, sex hormones (including steroids), anti-allergic agents, stimulants and anoretics, sympathomimetics, thyroid agents, vasodilators and xanthines. Preferred drug substances include those intended for oral administration and intravenous administration. A description of these classes of drugs and a listing of species within each class can be found in Martindale, *The Extra Pharmacopoeia*, Twenty-ninth Edition, The Pharmaceutical Press, London, 1989. The drug substances are commercially available and/or can be prepared by

45 techniques known in the art.

[0022] Representative illustrative species of drug substances useful in the practice of this invention include:

17- $\alpha$ -pregno-2,4-dien-20-yno-[2,3-d]-isoxazol-17-ol (danazol);  
 5 $\alpha$ ,17 $\alpha$ ,-1'-(methylsulfonyl)-1'H-pregn-20-yno [3,2-c]-pyrazol-17-ol (Steroid A);  
 50 [6-methoxy-4-(1-methylethyl)-3-oxo-1,2-benzisothiazol-2(3H)-yl]methyl 2,6-dichlorobenzoate 1,1-dioxide (WIN 63,394);  
 3-amino-1,2,4-benzotriazine-1,4-dioxide (WIN 59,075);  
 pivosulfam; pivosulfan; camptothecin; acetaminophen; acetylsalicylic acid; amiodarone; cholestyramine;  
 colestipol; cromolyn sodium; albuterol; sucralfat; sulfasalazine; minoxidil; tempazepam; alprazolam; propoxy-  
 55 phene; auranofin; erythromycin; cyclosporine; acyclovir; ganciclovir; etoposide; melphalan; methotrexate; mitoxantrone; daunorubicin; doxorubicin; megestrol; tamoxifen; medroxyprogesterone; nystatin; terbutaline; amphotericin B; aspirin; ibuprofen; naproxen; indomethacin; diclofenac; ketoprofen; flubiprofen; diflunisal; ethyl-3,5-diacetoamido-2,4,6-triiodobenzoate (WIN 8883);

ethyl-(3,5-bis(acetylamino)-2,4,6-triiodobenzoyloxy)acetate (WIN 12,901); and  
ethyl-2-(3,5-bis(acetylamino)-2,4,6-triiodobenzoyloxy)acetate (WIN 16,318).

5 [0023] In preferred embodiments of the invention, the drug substance is a steroid such as Danazol or Steroid A, an antiviral agent, an anti-inflammatory agent, an antineoplastic agent, a radiopharmaceutical or a diagnostic imaging agent.

[0024] The particles of this invention contain a discrete phase of a drug substance as described above having a surface modifier adsorbed on the surface thereof. Useful surface modifiers are believed to include those which physically adhere to the surface of the drug substance but do not chemically bond to the drug.

10 [0025] Suitable surface modifiers can preferably be selected from known organic and inorganic pharmaceutical excipients. Such excipients include various polymers, low molecular weight oligomers, natural products and surfactants. Preferred surface modifiers include nonionic and anionic surfactants. Representative examples of excipients include gelatin, casein, lecithin (phosphatides), gum acacia, cholesterol, tragacanth, stearic acid, benzalkonium chloride, calcium stearate, glyceryl monostearate, cetostearyl alcohol, cetomacrogol emulsifying wax, sorbitan esters, polyoxyethylene alkyl ethers, e.g., macrogol ethers such as cetomacrogol 1000, polyoxyethylene castor oil derivatives, polyoxyethylene sorbitan fatty acid esters, e.g., the commercially available Tweens™, polyethylene glycols, polyoxyethylene stearates, colloidal silicon dioxide, phosphates, sodium dodecylsulfate, carboxymethylcellulose calcium, carboxymethylcellulose sodium, methylcellulose, hydroxyethylcellulose, hydroxypropylcellulose, hydroxypropylmethylcellulose phthalate, noncrystalline cellulose, magnesium aluminum silicate, triethanolamine, polyvinyl alcohol (PVA), and polyvinylpyrrolidone (PVP). Most of these excipients are described in detail in the *Handbook of Pharmaceutical Excipients*, published jointly by the American Pharmaceutical Association and The Pharmaceutical Society of Great Britain, the Pharmaceutical Press, 1986. The surface modifiers are commercially available and/or can be prepared by techniques known in the art. Two or more surface modifiers can be used in combination.

20 [0026] Particularly preferred surface modifiers include polyvinylpyrrolidone, tyloxapol, polaxomers, such as Pluronic™ F68 and F108, which are block copolymers of ethylene oxide and propylene oxide available from BASF, and poloxamines, such as Tetronic™ 908 (T908), which is a tetrafunctional block copolymer derived from sequential addition of ethylene oxide and propylene oxide to ethylenediamine available from BASF, dextran, lecithin, Aerosol OT™, which is a dioctyl ester of sodium sulfosuccinic acid, available from American Cyanamid, Duponol™ P, which is a sodium lauryl sulfate, available from DuPont, Triton™ X-200, which is an alkyl aryl polyether sulfonate, available from Rohm and Haas, Tween 20 and Tween 80, which are polyoxyethylene sorbitan fatty acid esters, available from ICI Specialty Chemicals, Carbowax™ 3350 and 934, which are polyethylene glycols available from Union Carbide, Crodesta™ F-110, which is a mixture of sucrose stearate and sucrose distearate, available from Croda Inc., Crodesta SL-40, which is available from Croda Inc., and SA90HCO, which is  $C_{18}H_{37}-CH_2(CON(CH_3)CH_2(CHOH)_4CH_2OH)_2$ . Surface modifiers which have found to be particularly useful include polyvinylpyrrolidone, Pluronic F-68, and lecithin.

35 [0027] The surface modifier is adsorbed on the surface of the drug substance in an amount sufficient to maintain an effective average particle size of less than 400 nm. The surface modifier does not chemically react with the drug substance or itself. Furthermore, the individually adsorbed molecules of the surface modifier are essentially free of intermolecular crosslinkages. The surface modifier is adhered to the surfaces of the drug particles as contrasted to loading drugs into liposomes or polymers as in the prior techniques acknowledged above.

40 [0028] As used herein, particle size refers to a number average particle size as measured by conventional particle size measuring techniques well known to those skilled in the art, such as sedimentation field flow fractionation, photon correlation spectroscopy, or disk centrifugation. By "an effective average particle size of less than 400 nm", it is meant a particle size distribution in which at least 90% of the particles have a particle size of less than 400 nm when measured by the above-noted techniques. In preferred embodiments of the invention, the effective average particle size is less than 250 nm. In some embodiments of the invention, an effective average particle size of less than 100 nm has been achieved. With reference to the effective average particle size, it is preferred that at least 95% by weight and, more preferably, at least 99% by weight of the particles have a particle size less than the effective average particle size, e.g., 400 nm. In particularly preferred embodiments, essentially all of the particles have a size less than 400 nm. In some embodiments, essentially all of the particles have a size less than 250 nm.

50 [0029] The particles of this invention can be prepared in a method comprising the steps of dispersing a drug substance having a particle size larger than that ultimately desired in accordance with the invention in a liquid dispersion medium and applying mechanical means in the presence of grinding media to reduce the particle size of the drug substance to an effective average particle size of less than about 400 nm. The particles can be reduced in size in the presence of a surface modifier. Alternatively, the particles can be contacted with a surface modifier after attrition.

55 [0030] A general procedure for preparing the particles of this invention is set forth below. The drug substance selected is obtained commercially and/or prepared by techniques known in the art in a conventional coarse form. It is preferred, but not essential, that the particle size of the coarse drug substance selected be less than about 100  $\mu$ m as determined by sieve analysis. If the coarse particle size of the drug substance is greater than about 100  $\mu$ m, then it is

preferred that the particles of the drug substance be reduced in size to less than 100  $\mu\text{m}$  using a conventional milling method such as airjet or fragmentation milling.

**[0031]** The coarse drug substance selected can then be added to a liquid medium in which it is essentially insoluble to form a premix. The concentration of the drug substance in the liquid medium can vary from about 0.1 - 60%, and preferably is from 5 - 30% (w/w). It is preferred, but not essential, that the surface modifier be present in the premix. The concentration of the surface modifier can vary from about 0.1 to about 90%, and preferably is 1 - 75%, more preferably 20-60%, by weight based on the total combined weight of the drug substance and surface modifier. The apparent viscosity of the premix suspension is preferably less than about 1000 centipoise.

**[0032]** The premix can be used directly by subjecting it to mechanical means to reduce the average particle size in the dispersion to less than 400 nm. It is preferred that the premix be used directly when a ball mill is used for attrition. Alternatively, the drug substance and, optionally, the surface modifier, can be dispersed in the liquid medium using suitable agitation, e.g., a roller mill or a Cowles type mixer, until a homogeneous dispersion is observed in which there are no large agglomerates visible to the naked eye. It is preferred that the premix be subjected to such a premilling dispersion step when a recirculating media mill is used for attrition.

**[0033]** The mechanical means applied to reduce the particle size of the drug substance conveniently can take the form of a dispersion mill. Suitable dispersion mills include a ball mill, an attritor mill, a vibratory mill, and media mills such as a sand mill and a bead mill. A media mill is preferred due to the relatively shorter milling time required to provide the intended result, i.e., the desired reduction in particle size. For media milling, the apparent viscosity of the premix preferably is from about 100 to about 1000 centipoise. For ball milling, the apparent viscosity of the premix preferably is from about 1 up to about 100 centipoise. Such ranges tend to afford an optimal balance between efficient particle fragmentation and media erosion.

**[0034]** The grinding media for the particle size reduction step can be selected from rigid media preferably spherical or particulate in form having an average size less than about 3 mm and, more preferably, less than about 1 mm. Such media desirably can provide the particles of the invention with shorter processing times and impart less wear to the milling equipment. The selection of material for the grinding media is not believed to be critical. We have found that zirconium oxide, such as 95% ZrO stabilized with magnesia, zirconium silicate, and glass grinding media provide particles having levels of contamination which are believed to be acceptable for the preparation of pharmaceutical compositions. However, other media, such as stainless steel, titania, alumina, and 95% ZrO stabilized with yttrium, are expected to be useful. Preferred media have a density greater than about 3 g/cm<sup>3</sup>.

**[0035]** The attrition time can vary widely and depends primarily upon the particular mechanical means and processing conditions selected. For ball mills, processing times of up to five days or longer may be required. On the other hand, processing times of less than 1 day (residence times of one minute up to several hours) have provided the desired results using a high shear media mill.

**[0036]** The particles must be reduced in size at a temperature which does not significantly degrade the drug substance. Processing temperatures of less than about 30 - 40°C are ordinarily preferred. If desired, the processing equipment can be cooled with conventional cooling equipment. The method is conveniently carried out under conditions of ambient temperature and at processing pressures which are safe and effective for the milling process. For example, ambient processing pressures are typical of ball mills, attritor mills and vibratory mills. Processing pressures up to about 20 psi (1.4 kg/cm<sup>2</sup>) are typical of media milling.

**[0037]** The surface modifier, if it was not present in the premix, must be added to the dispersion after attrition in an amount as described for the premix above. Thereafter, the dispersion can be mixed, e.g., by shaking vigorously. Optionally, the dispersion can be subjected to a sonication step, e.g., using an ultrasonic power supply. For example, the dispersion can be subjected to ultrasonic energy having a frequency of 20 - 80 kHz for a time of about 1 to 120 seconds.

**[0038]** The relative amount of drug substance and surface modifier can vary widely and the optimal amount of the surface modifier can depend, for example, upon the particular drug substance and surface modifier selected, the critical micelle concentration of the surface modifier if it forms micelles, etc. The surface modifier preferably is present in an amount of about 0.1-10 mg per square meter surface area of the drug substance. The surface modifier can be present in an amount of 0.1-90%, preferably 20-60% by weight based on the total weight of the dry particle.

**[0039]** As indicated by the following examples, not every combination of surface modifier and drug substance provides the desired results. Consequently, the applicants have developed a simple screening process whereby compatible surface modifiers and drug substances can be selected which provide stable dispersions of the desired particles. First, coarse particles of a selected drug substance of interest are dispersed in a liquid in which the drug is essentially insoluble, e.g., water at 5% (w/w) and milled for 60 minutes in a DYNO-MILL under the standard milling conditions which are set forth in Example 1 which follows. The milled material is then divided into aliquots and surface modifiers are added at concentrations of 2, 10 and 50% by weight based on the total combined weight of the drug substance and surface modifier. The dispersions are then sonicated (1 minute, 20 kHz) to disperse agglomerates and subjected to particle size analysis by examination under an optical microscope (1000 x magnification). If a stable dispersion is observed, then the process for preparing the particular drug substance surface modifier combination can be optimized in accord-

ance with the teachings above. By stable it is meant that the dispersion exhibits no flocculation or particle agglomeration visible to the naked eye at least 15 minutes, and preferably, at least two days or longer after preparation.

[0040] The resulting dispersion of this invention is stable and consists of the liquid dispersion medium and the above-described particles. The dispersion of surface modified drug nanoparticles can be spray coated onto sugar spheres or onto a pharmaceutical excipient in a fluid-bed spray coater by techniques well known in the art.

[0041] Pharmaceutical compositions according to this invention include the particles described above and a pharmaceutically acceptable carrier therefor. Suitable pharmaceutically acceptable carriers are well known to those skilled in the art. These include non-toxic physiologically acceptable carriers, adjuvants or vehicles for parenteral injection, for oral administration in solid or liquid form, for rectal administration, and the like. A method of treating a mammal in accordance with this invention comprises the step of administering to the mammal in need of treatment an effective amount of the above-described pharmaceutical composition. The selected dosage level of the drug substance for treatment is effective to obtain a desired therapeutic response for a particular composition and method of administration. The selected dosage level therefore, depends upon the particular drug substance, the desired therapeutic effect, on the route of administration, on the desired duration of treatment and other factors. As noted, it is a particularly advantageous feature that the pharmaceutical compositions of this invention exhibit unexpectedly high bioavailability as illustrated in the examples which follow. Furthermore, it is contemplated that the drug particles of this invention provide more rapid onset of drug action in oral applications and decreased gastric irritancy.

[0042] It is contemplated that the pharmaceutical compositions of this invention will be particularly useful in oral and parenteral, including intravenous, administration applications. It is expected that poorly water soluble drug substances, which prior to this invention, could not have been administered intravenously, may be administered safely in accordance with this invention. Additionally, drug substances which could not have been administered orally due to poor bioavailability may be effectively administered in accordance with this invention.

[0043] While applicants do not wish to be bound by theoretical mechanisms, it is believed that the surface modifier hinders the flocculation and/or agglomeration of the particles by functioning as a mechanical or steric barrier between the particles, minimizing the close, interparticle approach necessary for agglomeration and flocculation. Alternatively, if the surface modifier has ionic groups, stabilization by electrostatic repulsion may result. It was surprising that stable drug particles of such a small effective average particle size and free of unacceptable contamination could be prepared by the method of this invention.

[0044] The following examples further illustrate the invention.

#### Example 1 - PVP Modified danazol particles prepared in a ball mill

[0045] A nanoparticulate dispersion of Danazol was prepared using a DYNO-MILL (Model KDL, manufactured by Willy A. Bachoffen AG Maschinenfabrik).

[0046] The following ingredients were added to a glass vessel and agitated on a roller for 24 hours to dissolve the polyvinylpyrrolidone surface modifier.

Polyvinylpyrrolidone K-15 (made by GAF) - 98 g  
High purity water - 664 g

[0047] Subsequently, 327 grams of dry powdered danazol was added to the above solution and rolled for one week. This step aided in evenly dispersing the Danazol in the surface modifier solution, thereby reducing the treatment time required in the media mill.

[0048] The danazol was purchased in a micronized form (average particle size of about 10 microns) from Sterling Winthrop Inc. The particles had been prepared by a conventional airjet milling technique.

[0049] This premix was added to a holding vessel and agitated with a conventional propeller mixer at low speed to maintain a homogeneous mixture for the media milling event. The media mill was prepared accordingly for the media milling process. The mill grinding chamber was partially filled with silica glass spheres and the premix was continuously recirculated through the media mill operating at the following conditions:

Grinding vessel: water jacketed stainless steel chamber  
Premix flow rate: 250 ml per minute  
Available volume of grinding vessel: 555 ml  
Media volume: 472 ml of glass beads  
Media type: size range of 0.5 - 0.75 mm silica glass beads, unleaded (distributed by Glen Mills, Inc.)  
Recirculation time: 240 min  
Residence time: 60 min  
Impeller speed: 3000 RPM, tangential speed 1952 ft/min



(595 m/min)

Grinding vessel coolant: water

Coolant temperature: 50°F (10°C)

- 5 [0050] After recirculating the slurry for 240 minutes, a sample of the dispersion was removed and evaluated for particle size distribution using a sedimentation field flow fractionator (made by DuPont). The particles were determined to have a number average diameter of 77.5 nm and a weight average diameter of 139.6 nm. The particle size of the dispersion ranged in size from 3 - 320 nm.

10 Example 2 - PVP modified danazol particles prepared in a ball mill at low solids.

[0051] A nanoparticulate dispersion of danazol was prepared using a ball mill process. A 600 ml cylindrical glass vessel (inside diameter = 3.0 inches (7.6 cm)) was filled approximately halfway with the following grinding media:

- 15 Grinding media: zirconium oxide grinding spheres (made by Zircoa, Inc.)

Media size: 0.85 - 1.18 mm diameter

Media volume: 300 ml

The following dry ingredients were added directly to this glass vessel:

danazol (micronized): 10.8 g

- 20 Polyvinylpyrrolidone K-15: 3.24 g

High purity water: 201.96 g

[0052] Danazol was purchased in the micronized form (average particle size 10 microns) from Sterling Winthrop Inc. and the polyvinylpyrrolidone was K-15 grade produced by GAF.

- 25 [0053] The cylindrical vessel was rotated horizontally about its axis at 57% of the "critical speed". The critical speed is defined as the rotational speed of the grinding vessel when centrifuging of the grinding media occurs. At this speed the centrifugal force acting on the grinding spheres presses and holds them firmly against the inner wall of the vessel. Conditions that lead to unwanted centrifuging can be computed from simple physical principles.

- 30 [0054] After 5 days of ball milling, the slurry was separated from the grinding media through a screen and evaluated for particle size with the sedimentation field flow fractionator. The number average particle diameter measured was 84.9 nm and the weight average particle diameter was 169.1 nm. The particles varied in size from 26 to 340 nm. The amount and type of surface modifier was sufficient to provide colloidal stability to agglomeration and to maintain a homogeneous blend of ingredients assuring precise material delivery during subsequent processing steps.

35 BIOAVAILABILITY TESTING

- [0055] Bioavailability of danazol from the nanoparticulate dispersion described above was compared to that from a suspension of unmilled danazol in fasted male beagle dogs. The unmilled material was prepared as a suspension in the same manner as the dispersion, with the exception of the ball milling process. Both formulations were administered to 40 each of five dogs by oral gavage and plasma obtained via a cannula in the cephalic vein. Plasma danazol levels were monitored over 24 hours. The relative bioavailability of danazol from the nanoparticulate dispersion was 15.9 fold higher than from the danazol suspension containing danazol particles having an average particle size of about 10 microns prepared by conventional airjet milling. Comparison of oral plasma levels with dose corrected plasma levels following intravenous administration of danazol gave a mean absolute bioavailability ( $\pm$  SEM) of  $82.3 \pm 10.1\%$  for the nanoparticulate 45 dispersion and  $5.1 \pm 1.9\%$  for the unmilled material.

Example 3 - PVP modified danazol particles prepared in a ball mill at high solids

- 50 [0056] A nanoparticle dispersion of danazol was prepared using 1 mm diameter glass grinding media (.85 - 1.18 mm from Potters Industries). A cylindrical glass vessel having a diameter of 2.75 inches (7.0 cm) with a volume of 400 ml was charged with 212 ml of unleaded glass grinding media. The following ingredients were added to this vessel:

30.4 g of micronized danazol

9.12 g of Polyvinylpyrrolidone K-15

- 55 112.48 g of high purity water

[0057] This vessel was rotated horizontally on its axis at a controlled rotational speed of 80.4 revolutions per minute (50% of critical speed) for 5 days. The slurry was immediately separated from the grinding media and evaluated for par-

particle size and grinding media attrition using inductively coupled plasma emissions (ICP). The particle size measured with a sedimentation field flow fractionator yielded a number average diameter of 112.7 nm and a weight average diameter of 179.3 nm. The extent of media attrition was measured to establish the purity of the final dispersion using an inductively coupled plasma-atomic emission spectroscopy method. The level of silicon in the final dispersion was less than 10 parts of elemental silicon per million parts of the slurry.

#### Example 4 - PVP modified danazol particles

**[0058]** A nanoparticle dispersion of danazol was prepared for clinical evaluation using a ball milling dispersion method. This dispersion was prepared by milling with glass grinding media. The grinding media used was:

Media type: 0.85 - 1.18 mm unleaded glass spheres

Media quantity: 6100 ml

The media was added to a 3 gallon porcelain jar. The following ingredients were then added to the jar:

1000 g danazol (micronized)

300 g Polyvinylpyrrolidone K-15

3700 g high purity water

**[0059]** The vessel was rolled 5 days at a rotational speed of 39.5 revolutions per minute (50% critical speed). The liquid slurry was separated from the grinding media with a screen and used to prepare solid oral doses for clinical studies. The dispersion was assessed for particle size using the sedimentation field flow fractionator and was measured to have a number average diameter of 134.9 nm and a weight average diameter of 222.2 nm. The level of contamination from the grinding media was measured (by ICP) to be 36 parts of silicon per million parts of dispersion. Less than 5 ppm of aluminum was detected. X-ray powder diffraction data of the starting powder was compared with the dispersed danazol and showed the crystal structure morphology of the solid dispersed particles was unchanged by the dispersion process.

#### Example 5 - PVP modified danazol particles

**[0060]** A nanoparticulate dispersion of danazol was prepared using a laboratory media mill and glass grinding media. The media mill was equipped with a 50 ml grinding chamber and the mill was a "Mini" Motormill manufactured by Eiger Machinery Inc.

**[0061]** The media mill was operated at the following process conditions:

Bead charge: 42.5 ml glass spheres

Rotor speed: 5000 RPM (2617 feet per minute (798 m/min) tangential speed)

Grinding media: 0.75 - 1.0 mm unleaded glass beads (distributed by Glens Mills)

**[0062]** The dispersion formula was prepared by dissolving 27 g of polyvinylpyrrolidone in 183 g of water and agitated in a steel vessel with a 50 mm "Cowles" type blade until the solution was clear and free of undissolved PVP polymer. The rotational speed of the mixer was maintained at 5000 RPM. 90 g of micronized danazol was slowly added to this blend with the same mixing for 30 min. 200 cc of the premix was added to the holding tank of the mill and recirculated for 5 hours and 51 minutes. The final residence time in the grinding zone was 40 minutes.

**[0063]** The final average particle size was measured and determined to have a number average diameter of 79.9 nm and a weight average diameter of 161.2 nm. The particles varied in size from 30 - 415 nm. The level of attrition from erosion of the grinding media and grinding vessel were measured (by ICP) to be 170 ppm of iron and 71 ppm silicon. The crystal structure was determined by x-ray diffraction to be unchanged by the dispersion process.

#### Example 6 - Lecithin modified Steroid A particles

**[0064]** A nanoparticulate dispersion of Steroid A was prepared by ball milling with zirconium oxide grinding beads. The dispersion was prepared in the absence of a surface modifier and a post addition of Lecithin and a sonication step were required to stabilize the dispersed phase of Steroid A and prevent agglomeration and rapid sedimentation.

**[0065]** A fine particle dispersion of Steroid A was prepared by ball milling the following ingredients:

5 g Steroid A

95 g high purity water

[0066] Steroid A was in the form of unmilled coarse grains having a particle size of about 100  $\mu\text{m}$  and ranging in size up to 400  $\mu\text{m}$ .

[0067] The following process conditions were used:

- 5 Media: 135 ml
- Vessel volume: 240 ml
- Media type: 0.85 - 1.18 mm Zirbeads (manufactured by Zircoa Inc.)
- Milling time: 4 days
- Milling speed: 86 RPM (50% critical speed)

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[0068] After four days of ball milling the slurry was separated from the grinding media through a screen. One gram of this unstabilized slurry was added to 10 g of an aqueous solution of Lecithin (1% Centrolex "P" by weight in high purity water, Lecithin manufactured by Central Soya Company, Inc.) and mixed by vigorous shaking, followed by a sonication step for 20 seconds using an ultrasonic horn (Model 350 Branson Ultrasonic Power Supply, Horn Diameter = 0.5

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inch (1.27 cm), Power setting = 2). The slurry was sized under a microscope. An Olympus BH-2 optical microscope equipped with phase contrast illumination was used to observe the size and condition of the dispersion.

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[0069] A drop of the above dilute slurry was placed between a microscope slide and glass cover slip and observed microscopically at high magnification (1,000 times) and compared to the slurry similarly diluted with water only (no surface modifier). The unmodified dispersion exhibited extensive particle agglomeration. The particle size of the unmodified dispersion was more than 10 microns and the unmodified dispersion exhibited no Brownian Motion. Brownian motion is the oscillatory or jiggling motion exhibited by particles in a liquid that fall in the size range of less than about 1 micron. The Lecithin modified particles exhibited rapid Brownian motion. The thus observed dispersion had the characteristics and appearance consistent with a number average particle size of less than 400 nm. Furthermore, it is expected that additional milling would lead to further particle size reduction so as to achieve an effective average particle size of less

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#### Example 7 - Alkyl aryl polyether sulfonate modified Steroid A

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[0070] Example 6 was repeated except that the Lecithin was replaced with Triton X-200 (manufacture by Rohm and Haas). Similar results were observed.

#### Example 8 - Gum acacia modified Steroid A

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[0071] Example 6 was repeated except that the Lecithin was replaced with gum acacia (available from Eastman Kodak Co.) Similar results were observed.

#### Example 9 - Sodium lauryl sulfate modified Steroid A

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[0072] Example 6 was repeated except that the Lecithin was replaced with sodium lauryl sulfate (available as Duponol ME from DuPont, Inc.). Similar results were observed.

#### Example 10 - Steroid A modified with a dioctylester of sodium sulfosuccinic acid

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[0073] Example 6 was repeated except that the Lecithin was replaced with Aerosol OT (available from American Cyanamid Chemical Products, Inc.). Similar results were observed.

#### Example 11 - Steroid A modified with a block copolymer of ethylene oxide and propylene oxide

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[0074] Example 6 was repeated except that the Lecithin was replaced with Pluronic F68 (available from BASF Corp.). Similar results were observed.

#### Example 12 - Steroid A particles modified with a block copolymer of ethylene oxide and propylene oxide

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[0075] A nanoparticulate dispersion of Steroid A was prepared by a ball milling process. The following ingredients were added to a cylindrical 0.95 l vessel. The vessel was filled approximately halfway with the following grinding media:

Grinding media: 0.85 - 1.18 mm diameter zirconium oxide spheres (made by Zircoa)

The following dispersion ingredients were added directly to the glass vessel:

18 g Steroid A  
 4.5 g Pluronic F68 (purchased from BASF Corp.)  
 336.6 g high purity water

5 [0076] Steroid A was purchased from Sterling Winthrop Inc. in the form of unmilled tabular crystals having an average particle size of approximately 100  $\mu\text{m}$ .

[0077] The vessel was rotated concentrically on its axis at 50% critical speed for 5 days. After this time 4.45 g of Pluronic F68 was added to the slurry and rolled for 5 more days at the same conditions. The slurry was then discharged and separated from the grinding media and evaluated for particle size using the sedimentation field flow fractionator.  
 10 The number average particle size measured was 204.6 nm and the weight average particle size was 310.6 nm. The particle size distribution ranged from approximately 68 - 520 nm. The dispersion was examined with an optical microscope. It exhibited excellent particle integrity, free of flocculation and agglomeration. The dispersion particles exhibited rapid Brownian motion.

#### 15 BIOAVAILABILITY TESTING

[0078] Bioavailability of Steroid A from the nanoparticulate dispersion described above was compared to that from a suspension of unmilled Steroid A (having an average particle size of about 100  $\mu\text{m}$ ) in male beagle dogs. The unmilled material was prepared as a suspension in the same manner as the dispersion, with the exception of the ball  
 20 milling process. Both formulations were administered to each of five dogs by oral gavage and plasma obtained via a cannula in the cephalic vein. Plasma Steroid A levels were monitored over 24 hours. The relative bioavailability of Steroid A from the nanoparticulate dispersion was 7.1 fold higher than from the unmilled Steroid A suspension. Comparison of oral plasma levels with dose corrected plasma levels following intravenous administration of Steroid A gave a mean absolute bioavailability ( $\pm$  SEM) of  $14.8 \pm 3.5\%$  for the nanoparticulate dispersion and  $2.1 \pm 1.0\%$  for the unmilled material.  
 25

#### Comparative Example A

[0079] A dispersion of Steroid A was prepared using a ball milling process with zirconium oxide grinding beads. The dispersion was prepared in the absence of a surface modifier and a post-sonication step was used to minimize flocculation and reaggregation.

[0080] A fine particle dispersion was prepared by ball milling the following ingredients:

5 g Steroid A  
 35 95 g high purity water  
 The following process conditions were used:  
 Grinding media: 135 ml  
 Vessel volume: 240 ml  
 Grinding media: 0.85 - 1.18 mm Zirbeads XR  
 40 Milling time: 4 days  
 Milling speed: 86 RPM (50% critical speed)

[0081] After four days of ball milling, the slurry was separated from the grinding media through a screen. One gram of the unstabilized slurry was blended with 10 grams of high purity water and mixed by vigorous shaking, followed by a  
 45 sonication step for 20 seconds using an ultrasonic horn (Model 350 Branson Ultrasonic Power Supply, Horn diameter = 0.5 inch, Power setting = 2). The slurry was sized under a microscope. An optical microscope equipped with phase contrast illumination was used to observe the condition of the dispersion.

[0082] A drop of the dilute slurry was placed between a microscope slide and a glass cover slip and observed at high magnification (400X). The dispersion exhibited severe particle aggregation. The aggregate size was greater than  
 50 10 microns and exhibited no Brownian particle movement.

#### Example 13-20

[0083] Table 1 is a summary of additional examples of the invention. Each of the examples in Table 1 resulted in  
 55 particles having an effective average particle size of less than 400 nm.

TABLE 1

Example	Class	Drug Substance	Surface Modifier	Number Average Particle Size
13.	anti-inflammatory	5% indomethacin	1% PVP	216 nm
14.	anti-inflammatory	5% indomethacin	1% F108	235 nm
15.	diagnostic imaging agent	10% WIN 8883	2% T908	166 nm
16.	diagnostic imaging agent	20% WIN 8883	3.3% T908	180 nm
17.	diagnostic imaging agent	20% WIN 8883	3.3% T908 (isotonic phosphate buffered saline, pH=7.4)	159 nm
18.	diagnostic imaging agent	20% WIN 8883	3.3% T908 (0.1M phosphate buffer pH=7.5)	167 nm
19.	diagnostic imaging agent	10% WIN 8883	1% SA9OHCO 1% Tween 20	194 nm
20.	diagnostic imaging agent	20% WIN 8883	1% SA9OHCO	193 nm

[0084] These examples demonstrate that the wet grinding process of this invention is broadly applicable to a wide variety of classes of poorly-soluble drug substances including steroids, anti-inflammatory agents, antineoplastic agents, radiopharmaceutical agents and diagnostic imaging agents having radically different chemical structures. Additionally, these examples demonstrate that the invention can be practiced in conjunction with a variety of surface modifiers and at a variety of surface modifier concentrations.

[0085] Furthermore, laboratory work has demonstrated that particles prepared according to this invention have exhibited a variety of unexpected properties, particularly with respect to increased bioavailability. For example, as described above, pharmaceutical compositions containing Steroid A and Danazol according to this invention have unexpectedly exhibited 7 and 16 fold increases in bioavailability compared to dispersions prepared by conventional techniques. Aqueous dispersions of WIN 63,394 prepared according to this invention resulted in an increase in bioavailability of 37-fold when compared to a conventional dispersion of WIN 63,394. The dispersions were administered at a dose of 5mg WIN 63,394 per kilogram of body weight to three dogs in the fasted state as a two way crossover study. Serial blood samples were withdrawn and analyzed by HPLC for WIN 63,394 concentrations. The relative bioavailabilities were calculated from the area under the curve for concentration versus time plots. Such increased bioavailability is particularly advantageous inasmuch as drug substances in the form of the particles of the instant invention can achieve the same therapeutic effect as substantially greater dosages of drug substances prepared by prior art techniques.

[0086] In addition, pharmaceutical compositions containing particles of this invention have exhibited improved dose proportionality and decreased fed-fasted variability. Further, particles of the invention comprising naproxen or indomethacin, when administered orally, have resulted in more rapid onset of action compared to conventional naproxen and indomethacin formulations. Moreover, certain of the particles of the invention have been found to be extraordinarily useful in x-ray contrast compositions.

#### Claims

1. Mechanically obtained particles having an effective average particle size of less than 400 nm, and being free of solvent contamination deriving from solvent precipitation, wherein the particles consist essentially of a crystalline drug substance and a surface modifier, said crystalline drug substance having been mechanically ground to an effective average particle size of less than 400 nm and having said surface modifier adhered to the surfaces of said particle essentially by adsorption with individually adsorbed molecules of said surface modifier being essentially free of intermolecular crosslinkages, said surface modifier being present in an amount of 0.1 to 90% by weight based on the total weight of dry particles so as to maintain said effective average particle size, wherein at least 90% of the particles have a weight average particle size of less than 400 nm, and wherein the surface modifier is selected so as to be compatible with the drug substance through a screening process so that the dispersion containing the particles exhibits no flocculation or particle agglomeration visible to the

naked eye and particularly when viewed under the optical microscope at 1000x at least two days after preparation.

2. Particles according to claim 1, in which the effective average particle size is less than about 250 nm.
- 5 3. Particles according to claim 2, in which the effective average particle size is less than about 100 nm.
4. Particles according to any one of the preceding claims, wherein said drug substance is selected from analgesics, anti-inflammatory agents, anthelmintics, anti-arrhythmic agents, antibiotics, anticoagulants, antidepressants, anti-diabetic agents, antiepileptics, antihistamines, antihypertensive agents, antimuscarinic agents, antimycobacterial agents, antineoplastic agents, immunosuppressants, antithyroid agents, antiviral agents, anxiolytic sedatives, astringents, beta-adrenoceptor blocking agents, blood products and substitutes, cardiac suppressants agents, contrast media, corticosteroids, cough suppressants, diagnostic agents, diagnostic imaging agents, diuretics, dopaminergics, haemostatics, immunological agents, lipid regulating agents, muscle relaxants, parasympathomimetics, parathyroid calcitonin and biphosphonates, prostaglandins, radio-pharmaceuticals, sex hormones, steroids, anti-allergic agents, stimulants and anoretics, sympathomimetics, thyroid agents, vasodilators and xanthines.
- 10 5. Particles according to any one of claims 1-3, wherein said drug substance is a steroid.
6. Particles according to any one of claims 1-3, wherein said drug substance is selected from the group consisting of danazol, 5 $\alpha$ ,17 $\alpha$ -1'-(methylsulfonyl)-1 H-pregn-20-yno[3,2-c]-pyrazol-17-ol, piposulfam, piposulfan, camptothecin, and ethyl-3,5-diacetamido-2,4,6-triiodobenzoate.
- 20 7. Particles according to any one of claims 1-3, wherein the drug substance is acetaminophen, acetylsalicylic acid, aspirin, ibuprofen, naproxen, indomethacin, diclofenac, ketoprofen, flubiprofen or diflunisal.
- 25 8. Particles according to any one of claims 1-3, wherein said drug substance is etoposide, piposulfam, piposulfan, camptothecin, melphalan, methotrexate, mitoxantrone, daunorubin, doxorubicin, or tamoxifen.
9. Particles according to any one of the claims 1-3, wherein said drug substance is cyclosporine, acyclovir, ganciclovir, medroxyprogesterone, or nystatin.
- 30 10. Particles according to any one of the preceding claims, wherein said surface modifier is gelatin, casein, lecithin, gum acacia, cholesterol, tragacanth, stearic acid, benzalkonium chloride, calcium stearate, glyceryl monostearate, cetostearyl alcohol, cetomacrogol emulsifying wax, a sorbitan ester, polyoxyethylene alkyl ethers, a polyoxyethylene castor oil derivative, a polyoxyethylene sorbitan fatty acid ester, a polyethylene glycol, a polyoxyethylene stearate, colloidal silicon dioxide, a phosphate, sodium dodecylsulfate, carboxymethylcellulose calcium, carboxymethylcellulose sodium, methylcellulose, hydroxyethylcellulose, hydroxypropylcellulose, hydroxypropylmethylcellulose phthalate, noncrystalline cellulose, magnesium aluminum silicate, triethanolamine, polyvinyl alcohol, polyvinylpyrrolidone, an ethylene oxide-propylene oxide block copolymer, an alkyl aryl polyether sulfonate, or a dioctylester of sodium sulfosuccinic acid.
- 40 11. Particles according to any one of claims 1-9, wherein said surface modifier is polyvinylpyrrolidone, tyloxapol, poloxamera, poloxamine, dextran, lecithin, a dioctylester of sodium sulfosuccinic acid, sodium lauryl sulfate, polyoxyethylene sorbitan, a fatty acid ester, a polyethylene glycol, or a mixture of sucrose stearate and sucrose distearate.
- 45 12. Particles according to any of claims 1-9, wherein said surface modifier is polyvinylpyrrolidone, tyloxapol, a poloxamer, a poloxamine, a polyoxyethylene sorbitan fatty acid ester, polyvinyl alcohol, hydroxy propyl cellulose or hydroxypropyl methyl cellulose.
- 50 13. Particles according to any of claims 1-9, wherein said surface modifier is polyvinylpyrrolidone, tyloxapol, poloxamer F108, a poloxamine, polyoxyethylene 20 sorbitan monolaurate, polyvinyl alcohol, hydroxy propyl cellulose or hydroxypropyl methyl cellulose.
- 55 14. Particles according to any one of claims 1-3, wherein said drug substance is crystalline 5 $\delta$ -17 $\delta$ , -1'-(methylsulfonyl) - 1' H-pregn-20-yno-pyrazol-17-ol ethylene oxide propylene oxide block copolymer.
15. A composition according to any one of the preceding claims, in which the drug substance has a solubility in water

of less than 10 mg./ml.

16. A stable dispersion consisting essentially of a liquid dispersion medium and particles according to any one of the preceding claims.
17. A dispersion according to claim 15, wherein said dispersion medium is water.
18. A dispersion according to claim 15, wherein said dispersion medium is safflower oil, ethanol, t-butanol, hexane or a glycol.
19. A pharmaceutical composition comprising particles according to any one of claims 1-14 and a pharmaceutically acceptable carrier therefor.
20. A method of preparing particles according to any one of claims 1-15, comprising the steps of dispersing larger size particles of the drug substance in a liquid dispersion medium and wet grinding said drug substance in the presence of rigid grinding media having an average particle size of less than 3 mm and the surface modifier to reduce the particle size of said drug substance to an effective average particle size of less than 400 nm.
21. A method of preparing particles according to any one of claims 1-15, comprising the steps of dispersing larger particles of the drug substance in a liquid dispersion medium, wet grinding said drug substance in the presence of rigid grinding media having an average particle size of less than 3 mm, and thereafter contacting said drug substance with a surface modifier by mixing said surface modifier with said dispersion medium to form particles having an effective average particle size of less than 400 nm.
22. A method according to claim 21, which includes subjecting the dispersion medium containing said drug substance and said surface modifier to ultrasonic energy.
23. A method according to any one of claims 20-22, wherein said grinding media have a density greater than 3 g/cm<sup>3</sup>.
24. A method according to any one of claims 20-23 wherein said grinding media have an average particle size of less than 1 mm.
25. A method according to any one of claims 20-24, in which the larger particle size drug is in the coarse form and which includes separating the ground particles from the grinding media.
26. Use of particles according to any one of claims 1-15, a dispersion according to any one of claims 16-18, a composition according to claim 19 or particles prepared according to any one of claims 20-25 for preparing a medicament.
27. Use according to claim 26, in which the medicament hastens the onset of action following administration.

#### Patentansprüche

1. Mechanisch erzeugte Partikeln, die eine effektive mittlere Korngröße von weniger als 400nm haben und die frei sind von Lösungsmittelverunreinigung durch die Lösungsmittelausfällung,  
wobei die Partikeln im wesentlichen aus einem kristallinen Arzneistoff und einem oberflächenmodifizierenden Stoff bestehen, der kristalline Arzneistoff mechanisch zu einer effektiven mittleren Korngröße von weniger als 400nm gemahlen wurde und der oberflächenmodifizierende Stoff an die - Oberfläche der Partikeln gebunden wurde, im wesentlichen durch Adsorption mit einzeln adsorbierten Molekülen des oberflächenmodifizierenden Stoffes, die im wesentlichen frei von Intermolekularbindungen sind, wobei der oberflächenmodifizierende Stoff in einer Menge von 0,1 bis 90 Gewichtsprozent, basierend auf dem Gesamtgewicht der trockenen Partikeln, vorliegt, so daß die effektive mittlere Korngröße erhalten bleibt, wobei mindestens 90% der Partikeln eine gewichtete mittlere Korngröße von weniger als 400nm haben und wobei der oberflächenmodifizierende Stoff durch ein Screeningverfahren so gewählt ist, daß er sich mit dem Arzneistoff verträgt, damit die Dispersion, die die Partikeln enthält, weder Ausflockung noch Partikelagglomeration zeigt, die mit bloßem Auge und insbesondere bei Betrachtung unter dem optischen Mikroskop bei 1000facher Vergrößerung mindestens zwei Tage nach der Herstellung sichtbar sind.
2. Partikeln nach Anspruch 1, bei denen die effektive mittlere Korngröße weniger als etwa 250nm beträgt.

3. Partikeln nach Anspruch 2, bei denen die effektive mittlere Korngröße weniger als etwa 100nm beträgt.
4. Partikeln nach einem der vorigen Ansprüche, wobei der Arzneistoff aus Analgetika, antiphlogistisch wirkenden Mitteln, Anthelminthika, antiarrhythmisch wirkenden Mitteln, Antibiotika, Antikoagulantien, Antidepressiva, antidiabetisch wirkenden Mitteln, Antiepileptika, Antihistaminika, antihypertensiv wirkenden Mitteln, antimuskarinisch wirkenden Mitteln, antimycobakteriell wirkenden Mitteln, antineoplastisch wirkenden Mitteln, Immunsuppressiva, thyreostatisch wirkenden Mitteln, antiviral wirkenden Mitteln, anxiolytisch wirksamen Sedativa, Adstringentien, Betablockern, Blutprodukten und -ersatzmitteln, herzwirksamen Suppressiva, Kontrastmitteln, Corticosteroiden, Hustenstillern, diagnostischen Mitteln, diagnostischen Bildgebungsmitteln, Diuretika, Dopaminagonisten, blutstillenden Mitteln, immunologisch wirkenden Mitteln, lipidregulierenden Mitteln, Muskelrelaxantien, Parasympathomimetika, parathyreoidalem Calcitonin und Bisphosphonaten, Prostaglandinen, Radiopharmaka, Geschlechtshormonen, Steroiden, antiallergisch wirkenden Mitteln, Stimulantien und Anorektika, Sympathomimetika, auf die Schilddrüse wirkenden Mitteln, Vasodilatoren und Xanthinen gewählt ist.
5. Partikeln nach einem der Ansprüche 1-3, wobei der Arzneistoff ein Steroid ist.
6. Partikeln nach einem der Ansprüche 1-3, wobei der Arzneistoff aus der Gruppe bestehend aus Danazol, 5 $\alpha$ , 17 $\alpha$ -1'-(methylsulfonyl)-1 H-pregn-20-yno[3,2-c]-pyrazol-17-01, Pisosulfam, Pisosulfan, Camptothecin, und Ethyl-3,5-diacetamido-2,4,6-trijodobenzoat gewählt ist.
7. Partikeln nach einem der Ansprüche 1-3, wobei der Arzneistoff Acetaminophen, Acetylsalicylsäure, Aspirin, Ibuprofen, Naproxen, Indomethacin, Diclofenac, Ketoprofen, Flubiprofen oder Diflunisal ist.
8. Partikeln nach einem der Ansprüche 1-3, wobei der Arzneistoff Etoposid, Pisosulfam, Pisosulfan, Camptothecin, Melphalan, Methotrexat, Mitoxantron, Daunorubin, Doxorubicin, oder Tamoxifen ist.
9. Partikeln nach einem der Ansprüche 1-3, wobei der Arzneistoff Ciclosporin, Aciclovir, Gancilovar, Medroxyprogesteron oder Nystatin ist.
10. Partikeln nach einem der vorigen Ansprüche, wobei der oberflächenmodifizierende Stoff Gelatine, Casein, Lecithin, arabisches Gummi, Cholesterin, Tragant, Stearinsäure, Benzalkoniumchlorid, Calciumstearat, Glycerolmonostearat, Cetylstearylalkohol, emulgierendes Cetylmacrogol-Wachs, ein Sorbitanester, Polyoxyethylenalkylether, ein Polyoxyethylenrizinusölderivat, ein Polyoxyethylensorbitanfettsäureester, ein Polyethylenglycol, ein Polyoxyethylenstearat, kolloiddisperses Siliciumdioxid, ein Phosphat, Natriumdodecylsulfat, Carboxymethylcellulose-Calcium, Carboxymethylcellulose-Natrium, Methylcellulose, Hydroxyethylcellulose, Hydroxypropylcellulose, Hydroxypropylmethylcellulosephthalat, nichtkristalline Cellulose, Magnesiumaluminiumsilicat, Triethanolamin, Polyvinylalkohol, Polyvinylpyrrolidon, ein Ethylenoxidpropylenoxid-Blockcopolymer, ein Alkylarylpolyethersulfonat oder ein Dioctylester der Natriumsulfobornsteinsäure ist.
11. Partikeln nach einem der Ansprüche 1-9, wobei der oberflächenmodifizierende Stoff Polyvinylpyrrolidon, Tyloxapol, Poloxamer, Poloxamin, Dextran, Lecithin, ein Dioctylester der Natriumsulfobornsteinsäure, Natriumlaurylsulfat, Polyoxyethylensorbitan, ein Fettsäureester, ein Polyethylenglykol oder eine Mischung aus Saccharosestearat und Saccharosedistearat ist.
12. Partikeln nach einem der Ansprüche 1-9, wobei der oberflächenmodifizierende Stoff Polyvinylpyrrolidin, Tyloxapol, ein Poloxamer, ein Poloxamin, ein Polyoxyethylensorbitanfettsäureester, Polyvinylalkohol, Hydroxypropylcellulose oder Hydroxypropylmethylcellulose ist.
13. Partikeln nach einem der Ansprüche 1-9, wobei der oberflächenmodifizierende Stoff Polyvinylpyrrolidin, Tyloxapol, Poloxamer F 108, ein Poloxamin, Polyoxyethylen 20 sorbitanmonolaurat, Polyvinylalkohol, Hydroxypropylcellulose oder Hydroxypropylmethylcellulose ist.
14. Partikeln nach einem der Ansprüche 1-3, wobei der Arzneistoff kristallines 5 $\delta$ -17 $\delta$ , -1'-(Methylsulfonyl)-1' H-pregn-20-yno-pyrazol-17-ol Ethylenoxidpropylenoxid-Blockcopolymer ist.
15. Eine Zusammensetzung nach einem der vorigen Ansprüche, in der der Arzneistoff eine Wasserlöslichkeit von weniger als 10mg/ml besitzt.



16. Eine stabile Dispersion, die im wesentlichen aus einem flüssigen Dispersionsmittel und Partikeln nach einem der vorigen Ansprüche besteht.
17. Eine Dispersion nach Anspruch 15, wobei das Dispersionsmittel Wasser ist.
18. Eine Dispersion nach Anspruch 15, wobei das Dispersionsmittel Färberdistelöl, Ethanol, t-Butanol, Hexan oder ein Glykol ist.
19. Eine pharmazeutische Zusammensetzung, die Partikeln aus einem der Ansprüche 1-14 sowie einen damit pharmazeutisch verträglichen Träger enthält.
20. Ein Verfahren zur Herstellung von Partikeln nach einem der Ansprüche 1-15, das folgende Schritte umfaßt: Dispergieren von größeren Arzneistoffpartikeln in einem flüssigen Dispersionsmittel und das nasse Mahlen des Arzneistoffs in Gegenwart eines festen Mahlmediums, das eine mittlere Korngröße von weniger als 3mm besitzt, und des oberflächenmodifizierenden Stoffes, um die Korngröße des Arzneistoffs zu einer effektiven mittleren Korngröße von weniger als 400nm zu reduzieren.
21. Ein Verfahren zur Herstellung von Partikeln aus einem der Ansprüche 1-15, das folgende Schritte umfaßt: Dispergieren größerer Arzneistoffpartikeln in einem flüssigen Dispersionsmittel, das nasse Mahlen des Arzneistoffs in Gegenwart eines festen Mahlmediums, das eine mittlere Korngröße von weniger als 3mm besitzt, und danach das In-Kontakt-bringen des Arzneistoffs mit einem oberflächenmodifizierenden Stoff, indem der oberflächenmodifizierende Stoff mit dem Dispersionsmittel vermischt wird, um Partikeln zu bilden, die eine effektive mittlere Korngröße von weniger als 400nm besitzen.
22. Ein Verfahren nach Anspruch 21, das eine Behandlung des Dispersionsmittels mit dem darin enthaltenen Arzneistoff und dem oberflächenmodifizierenden Stoff mit Ultraschallenergie beinhaltet.
23. Ein Verfahren nach einem der Ansprüche 20-22, wobei das Mahlmedium eine größere Dichte als  $3\text{g/cm}^3$  besitzt.
24. Ein Verfahren nach einem der Ansprüche 20-23, wobei das Mahlmedium eine mittlere Korngröße von weniger als 1mm besitzt.
25. Ein Verfahren nach einem der Ansprüche 20-24, in dem der Arzneistoff mit größerer Korngröße in grobkörniger Form vorliegt und das die Trennung der gemahlten Partikeln vom Mahlmedium umfaßt.
26. Die Verwendung von Partikeln nach einem der Ansprüche 1-15, einer Dispersion nach einem der Ansprüche 16-18, einer Zusammensetzung nach Anspruch 19 oder Partikeln, die nach einem der Ansprüche 20-25 zur Herstellung eines Medikamentes hergestellt wurden.
27. Die Verwendung nach Anspruch 26, wobei das Medikament den Wirkungseintritt, der auf die Verabreichung folgt, beschleunigt.

#### Revendications

1. Particules obtenues mécaniquement ayant une granulométrie moyenne efficace inférieure à 400 nm, et étant dépourvues de contamination par un solvant dérivant d'une précipitation dans un solvant, dans lesquelles les particules sont essentiellement constituées d'une substance cristalline de type médicament et d'un agent de modification de surface, ladite substance cristalline de type médicament ayant été broyée mécaniquement à une granulométrie moyenne efficace inférieure à 400 nm et ledit agent de modification de surface étant collé aux surfaces desdites particules essentiellement par adsorption, les molécules adsorbées individuellement dudit agent de modification de surface étant essentiellement dépourvues de réticulations intermoléculaires, ledit agent de modification de surface étant présent en une proportion de 0,1 à 90 % en poids, par rapport au poids total des particules sèches de façon à conserver ladite granulométrie moyenne efficace, dans lesquelles au moins 90 % des particules ont une granulométrie moyenne en poids inférieure à 400 nm, et dans lesquelles l'agent de modification de surface est choisi de façon à être compatible avec la substance de type médicament lors d'un procédé de tamisage de sorte que la dispersion contenant les particules ne présente pas de floculation ni d'agglomération des particules visible à l'oeil nu et, en particulier, lorsqu'elles sont examinées au microscope optique sous un grossissement de 1000 au moins deux jours après la préparation.

2. Particules selon la revendication 1, dans lesquelles la granulométrie moyenne efficace est inférieure à environ 250 nm.
3. Particules selon la revendication 2, dans lesquelles la granulométrie moyenne efficace est inférieure à environ 100 nm.
4. Particules selon l'une quelconque des revendications précédentes, dans lesquelles ladite substance de type médicament est choisie parmi les analgésiques, les agents anti-inflammatoires, les anthelminthiques, les agents anti-arythmiques, les antibiotiques, les anticoagulants, les antidépresseurs, les agents antidiabétiques, les anti-épileptiques, les antihistaminiques, les agents antihypertenseurs, les agents antimuscariniques, les agents antimycobactériens, les agents antinéoplastiques, les immunosuppresseurs, les agents antithyroïdiens, les agents antiviraux, les sédatifs anxiolytiques, les astringents, les agents bêta-bloquants des adrénocéphes, les produits et les produits de substitution sanguins, les supprimeurs cardiaques, les agents de contraste, les corticostéroïdes, les anti-tussifs, les agents de diagnostic, les agents d'imagerie diagnostique, les diurétiques, les dopaminergiques, les hémostatiques, les agents immunologiques, les agents de régulation des lipides, les relaxants musculaires, les parasymphomimétiques, la calcitonine parathyroïdienne et les biphosphonates, les prostaglandines, les agents radio-pharmaceutiques, les hormones sexuelles, les stéroïdes, les agents anti-allergiques, les stimulants et les anorexiantes, les sympathomimétiques, les agents thyroïdiens, les vasodilatateurs et les xanthines.
5. Particules selon l'une quelconque des revendications 1 à 3, dans lesquelles ladite substance de type médicament est un stéroïde.
6. Particules selon l'une quelconque des revendications 1 à 3, dans lesquelles ladite substance de type médicament est choisie dans le groupe constitué par le danazol, le 5 $\alpha$ -17 $\alpha$ -1'-(méthylsulfonyl)-1-H-prégn-20-yno[3,2-c]-pyrazol-17-ol, la piposulfame, le piposulfane, la camptothécine et le 3,5-diacétamido-2,4,6-triiodobenzoate d'éthyle.
7. Particules selon l'une quelconque des revendications 1 à 3, dans lesquelles la substance de type médicament est l'acétaminophène, l'acide acétylsalicylique, l'aspirine, l'ibuprofène, le naproxène, l'indométhacine, le diclofénac, le cétoprofène, le flubiprofène ou le diflunisal.
8. Particules selon l'une quelconque des revendications 1 à 3, dans lesquelles ladite substance de type médicament est l'étoposide, la piposulfame, le piposulfane, la camptothécine, le melphalan, le méthotrexate, la mitoxantrone, la daunorubine, la doxorubicine ou le tamoxifène.
9. Particules selon l'une quelconque des revendications 1 à 3, dans lesquelles ladite substance de type médicament est la cyclosporine, l'acyclovir, le ganciclovir, la médroxyprogestérone ou la nystatine.
10. Particules selon l'une quelconque des revendications précédentes, dans lesquelles ledit agent de modification de surface est la gélatine, la caséine, la lécithine, la gomme arabique, le cholestérol, la gomme adragante, l'acide stéarique, le chlorure de benzalkonium, le stéarate de calcium, le monostéarate de glycéryle, l'alcool cétostéarylique, une cire émulsifiante de type céto-macrogol, un ester de sorbitane, les éthers alkylés polyoxyéthylés, un dérivé d'huile de ricin polyoxyéthylé, un ester d'acide gras de sorbitane polyoxyéthylé, un polyéthylène glycol, un stéarate polyoxyéthylé, le dioxyde de silicium colloïdal, un phosphate, le dodécylsulfate de sodium, la carboxyméthylcellulose de calcium, la carboxyméthylcellulose de sodium, la méthylcellulose, l'hydroxyéthylcellulose, l'hydroxypropylcellulose, le phtalate d'hydroxypropylméthylcellulose, la cellulose non cristalline, l'aluminosilicate de magnésium, la triéthanolamine, le poly(alcool vinylique), la polyvinylpyrrolidone, un copolymère séquencé oxyde d'éthylène-oxyde de propylène, un alkylarylpolysulfonate, ou un dioctylester d'acide sulfosuccinique de sodium.
11. Particules selon l'une quelconque des revendications 1 à 9, dans lesquelles ledit agent de modification de surface est la polyvinylpyrrolidone, le tyloxapol, un poloxamère, une poloxamine, le dextrane, la lécithine, un dioctylester d'acide sulfosuccinique de sodium, le laurylsulfate de sodium, le sorbitane polyoxyéthylé, un ester d'acide gras, un polyéthylène glycol, ou un mélange de stéarate de saccharose et de distéarate de saccharose.
12. Particules selon l'une quelconque des revendications 1 à 9, dans lesquelles ledit agent de modification de surface est la polyvinylpyrrolidone, le tyloxapol, un poloxamère, une poloxamine, un ester d'acide gras de sorbitane polyoxyéthylé, le poly(alcool vinylique), l'hydroxypropylcellulose ou l'hydroxypropylméthylcellulose.

13. Particules selon l'une quelconque des revendications 1 à 9, dans lesquelles ledit agent de modification de surface est la polyvinylpyrrolidone, le tyloxapol, le poloxamère F108, une poloxamine, le monolaurate de sorbitane polyoxyéthyléné (20), le poly(alcool vinylique), l'hydroxypropylcellulose ou l'hydroxypropylméthylcellulose.
- 5 14. Particules selon l'une quelconque des revendications 1 à 3, dans lesquelles ladite substance de type médicament est un copolymère séquencé cristallin 5 $\alpha$ -17 $\alpha$ -1'-(méthylsulfonyl)-1'-H-prégn-20-yno-pyrazol-17-ol/oxyde d'éthylène/oxyde de propylène.
- 10 15. Composition selon l'une quelconque des revendications précédentes, dans laquelle la substance de type médicament a une solubilité dans l'eau inférieure à 10 mg/ml.
16. Dispersion stable essentiellement constituée d'un milieu de dispersion liquide et des particules selon l'une quelconque des revendications précédentes.
- 15 17. Dispersion selon la revendication 15, dans laquelle ledit milieu de dispersion est l'eau.
18. Dispersion selon la revendication 15, dans laquelle ledit milieu de dispersion est l'huile de tournesol, l'éthanol, le t-butanol, l'hexane ou un glycol.
- 20 19. Composition pharmaceutique comprenant les particules selon l'une quelconque des revendications 1 à 14, et un véhicule pharmaceutiquement acceptable de celles-ci.
20. Procédé de préparation des particules selon l'une quelconque des revendications 1 à 15, comprenant les étapes consistant à disperser des particules plus grosses de la substance de type médicament dans un milieu de dispersion liquide et à broyer à l'état humide ladite substance de type médicament en présence de moyens de broyage rigides ayant une granulométrie moyenne inférieure à 3 mm et l'agent de modification de surface pour réduire la granulométrie de ladite substance de type médicament à une granulométrie moyenne efficace inférieure à 400 nm.
- 25 21. Procédé de préparation des particules selon l'une quelconque des revendications 1 à 15, comprenant les étapes consistant à disperser des particules plus grosses de la substance de type médicament dans un milieu de dispersion liquide, à broyer à l'état humide ladite substance de type médicament en présence de moyens de broyage rigides ayant une granulométrie moyenne inférieure à 3 mm, puis à mettre en contact ladite substance de type médicament avec un agent de modification de surface par mélange dudit agent de modification de surface avec ledit milieu de dispersion pour former des particules ayant une granulométrie moyenne efficace inférieure à 400 nm.
- 30 22. Procédé selon la revendication 21, qui comprend le fait de soumettre le milieu de dispersion contenant ladite substance de type médicament et ledit agent de modification de surface à une énergie ultrasonore.
- 40 23. Procédé selon l'une quelconque des revendications 20 à 22, dans lequel lesdits moyens de broyage ont une masse volumique supérieure à 3 g/cm<sup>3</sup>.
24. Procédé selon l'une quelconque des revendications 20 à 23, dans lequel lesdits moyens de broyage ont une granulométrie moyenne inférieure à 1 mm.
- 45 25. Procédé selon l'une quelconque des revendications 20 à 24, dans lequel le médicament de plus grande taille de particule est sous forme grossière et qui comprend la séparation des particules broyées des milieux de broyage.
- 50 26. Utilisation des particules selon l'une quelconque des revendications 1 à 15, d'une dispersion selon l'une quelconque des revendications 16 à 18, d'une composition selon la revendication 19 ou des particules préparées selon l'une quelconque des revendications 20 à 25, pour préparer un médicament.
27. Utilisation selon la revendication 26, dans laquelle le médicament accélère le démarrage de l'action après l'administration.

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